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(54) Title: IMIDAZOLE DERIVATIVES AND THEIR USE AS CYTOKINE INHIBITORS

(57) Abstract

Novel 2,4,5-triarylimidazole compounds and compositions for use in therapy.

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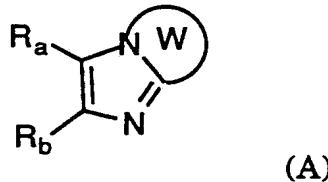
IMIDAZOLE DERIVATIVES AND THEIR USE AS CYTOKINE INHIBITORS

This invention relates to a novel group of imidazole compounds, processes for the preparation thereof, the use thereof in treating cytokine mediated diseases and pharmaceutical compositions for use in such therapy.

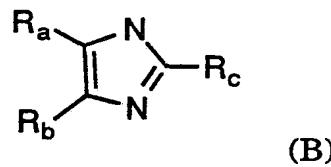
BACKGROUND OF THE INVENTION:

There has been much interest in the past few years in compounds which are cytokine-inhibitors, for use in treating disease states which are associated with the excessive or unregulated production of cytokines.

Compounds of the general formula (A):



wherein R_a is pyridyl, R_b is optionally substituted phenyl and W is a partially or fully unsaturated fused 5- or 6-membered heterocyclic ring, such as pyrrolyl, pyridyl, dihydropyrrolyl, dihydropyridinyl, dihydrothiazolyl or tetrahydro-triazinyl, are inhibitors of the cytokines IL-1 and TNF (see WO88/01169, WO90/15534, WO91/00092, WO92/10190, WO92/10498 and WO92/12154, published after the filing of this application). In addition, these compounds are also inhibitors of the enzyme 5-lipoxygenase. We have now surprisingly found that if the ring W is replaced by certain substituents at the 2-position, cytokine-inhibitory activity is maintained. Such compounds are generically 2-substituted-4-aryl-5-heteroaryl-imidazoles. Compounds within this class have already been extensively investigated, as anti-inflammatory agents, acting principally as cyclo-oxygenase inhibitors, as described in, for instance, US patents 3,707,405 and 3,929,807. The latter discloses compounds of the general formula (B):



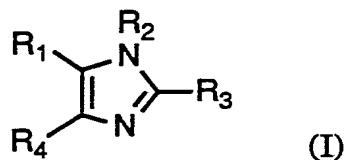
wherein one of R_a and R_b is optionally substituted phenyl and the other is a 6-membered heterocyclic ring with 1 or 2 nitrogen atoms and R_c represents lower alkyl, cycloalkyl or phenyl optionally substituted by halogen, lower alkyl or lower alkoxy, in particular the compound 2-(4-chlorophenyl)-4-(4-methoxyphenyl)-5-(4-pyridyl)-imidazole. These compounds are said to have

anti-inflammatory, analgesic and antipyretic activity. There is however no mention that these compounds may be cytokine inhibitors.

FULL DESCRIPTION OF THE INVENTION:

5 Accordingly, the present invention provides a compound of formula

(I):



wherein:

R₁ is 4-pyridyl, pyrimidinyl, quinolyl, isoquinolinyl, 1-imidazolyl or 1-

10 benzimidazolyl which is optionally substituted with one or two substituents each of which is independently selected from C₁-4 alkyl, halo, C₁-4 alkoxy, C₁-4 alkylthio, NH₂, mono- or di-C₁-6-alkylamino or N-heterocyclyl ring which ring has from 5 to 7 members and optionally contains an additional heteroatom selected from oxygen, sulfur or NR₂₂;

15 R₂ is R₈ or -OR₁₂;

R₃ is -X_aP(Z)(X_bR₁₃)₂ or Q-(Y₁)_t;

Q is an aryl or heteroaryl group;

t is an integer of 1 to 3;

X_a is -NR₈- , -O- , -S- or a C₁-10 alkylene chain optionally substituted by C₁-4 alkyl and optionally interrupted by -NR₈- , -O- or -S-;

X_b is -(CR₁₀R₂₀)_n , -NR₈- , -O- or -S-;

Z is oxygen or sulfur;

n is 0 or an integer from 1 to 10;

Y₁ is independently selected from hydrogen, C₁-5 alkyl, halo-substituted C₁-5 alkyl, halogen, -X_a-P(Z)-(X_bR₁₃)₂ or -(CR₁₀R₂₀)_nY₂;

Y₂ is -OR₈ , -NO₂ , -S(O)_mR₁₁ , -SR₈ , -S(O)_mOR₈ , -S(O)_mNR₈R₉ , -NR₈R₉ ,

-O(CR₁₀R₂₀)_nNR₈R₉ , -C(O)R₈ , -CO₂R₈ , -CO₂(CR₁₀R₂₀)_nCONR₈R₉ ,

-ZC(O)R₈ , -CN , -C(Z)NR₈R₉ , -NR₁₀C(Z)R₈ , -C(Z)NR₈OR₉ ,

-NR₁₀C(Z)NR₈R₉ , -NR₁₀S(O)_mR₁₁ , -N(OR₂₁)C(Z)NR₈R₉ , -N(OR₂₁)C(Z)R₈ ,

30 -C(=NOR₂₁)R₈ , -NR₁₀C(=NR₁₅)SR₁₁ , -NR₁₀C(=NR₁₅)NR₈R₉ ,

-NR₁₀C(=CR₁₄R₂₄)SR₁₁ , -NR₁₀C(=CR₁₄R₂₄)NR₈R₉ ,

-NR₁₀C(O)C(O)NR₈R₉ , -NR₁₀C(O)C(O)OR₁₀ , -C(=NR₁₃)NR₈R₉ ,

-C(=NOR₁₃)NR₈R₉ , -C(=NR₁₃)ZR₁₁ , -OC(Z)NR₈R₉ , -NR₁₀S(O)_mCF₃ ,

-NR₁₀C(Z)OR₁₀ , 5-(R₁₈)-1,2,4-oxadizaol-3-yl or 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-

35 dihydro-1,2,4-oxadiazol-3-yl;

m' is 1 or 2;

n' is an integer from 1 to 10;

R₄ is phenyl, naphth-1-yl or naphth-2-yl which is optionally substituted by one or two substituents, each of which is independently selected, and which, for a 4-phenyl, 4-naphth-1-yl or 5-naphth-2-yl substituent, is halo, cyano, -C(Z)NR₇R₁₇, -C(Z)OR₂₃, -(CR₁₀R₂₀)_mCOR₃₆, -SR₅, -SOR₅, -OR₃₆, 5 halo-substituted-C₁₋₄ alkyl, C₁₋₄ alkyl, -ZC(Z)R₃₆, -NR₁₀C(Z)R₂₃, or -(CR₁₀R₂₀)_mNR₁₀R₂₀ and which, for other positions of substitution, is halo, cyano, -C(Z)NR₁₆R₂₆, -C(Z)OR₈, -(CR₁₀R₂₀)_mCOR₈, -S(O)_mR₈, -OR₈, halo-substituted-C₁₋₄ alkyl, -C₁₋₄ alkyl, -(CR₁₀R₂₀)_mNR₁₀C(Z)R₈, 10 -NR₁₀S(O)_mR₁₁, -NR₁₀S(O)_mNR₇R₁₇ wherein m is 1 or 2, -ZC(Z)R₈ or -(CR₁₀R₂₀)_mNR₁₆R₂₆;

m is 0, or the integer 1 or 2;

R₅ is hydrogen, C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl or NR₇R₁₇, excluding the moieties -SR₅ being -SNR₇R₁₇ and -SOR₅ being -SOH;

R₆ is C₁₋₄ alkyl, halo-substituted-C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl or C₃₋₅ 15 cycloalkyl;

R₇ and R₁₇ is each independently selected from hydrogen or C₁₋₄ alkyl or R₇ and R₁₇ together with the nitrogen to which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or NR₂₂;

R₈ is hydrogen, heterocyclyl, heterocyclylalkyl or R₁₁;

R₉ is hydrogen, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, C₃₋₇ cycloalkyl, 20 C₅₋₇ cycloalkenyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl or R₈ and R₉ may together with the nitrogen to which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or NR₁₂;

R₁₀ and R₂₀ is each independently selected from hydrogen or C₁₋₄ alkyl;

R₁₁ is C₁₋₁₀ alkyl, halo-substituted C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ 25 alkynyl, C₃₋₇ cycloalkyl, C₅₋₇ cycloalkenyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl;

R₁₂ is hydrogen, -C(Z)R₁₃ or optionally substituted C₁₋₄ alkyl, optionally substituted aryl or optionally substituted aryl-C₁₋₄ alkyl;

R₁₃ is hydrogen, C₁₋₁₀ alkyl, cycloalkyl, heterocyclyl, aryl, arylalkyl, 30 heteroaryl or heteroarylalkyl;

R₁₄ and R₂₄ is each independently selected from hydrogen, alkyl, nitro or cyano; R₁₅ is hydrogen, cyano, C₁₋₄ alkyl, C₃₋₇ cycloalkyl or aryl;

R₁₆ and R₂₆ is each independently selected from hydrogen or optionally substituted C₁₋₄ alkyl, optionally substituted aryl or optionally substituted aryl-C₁₋₄ alkyl, or together with the nitrogen which they are 35

attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or NR₁₂ ; R₁₈ and R₁₉ is each independently selected from hydrogen, C₁₋₄ alkyl, substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl or together denote a oxygen or sulfur;

5 R₂₁ is hydrogen, a pharmaceutically acceptable cation, C₁₋₁₀ alkyl, C₃₋₇ cycloalkyl, aryl, aryl C₁₋₄ alkyl, heteroaryl, heteroarylalkyl, heterocyclyl, aroyl, or C₁₋₁₀ alkoyl ;

R₂₂ is R₁₀ or C(Z)-C₁₋₄ alkyl;

10 R₂₃ is C₁₋₄ alkyl, halo-substituted-C₁₋₄ alkyl, or C₃₋₅ cycloalkyl ; R₃₆ is hydrogen or R₂₃ ; or a pharmaceutically acceptable salt thereof ; and excluding 2-(4-chlorophenyl)-4-(4-methoxyphenyl)-5-(4-pyridyl)imidazole.

15 Suitable R₁ moieties include 4-pyridyl, 4-pyrimidinyl, 4-quinolyl, 6-isoquinolinyl, 1-imidazolyl and 1-benzimidazolyl, of which 4-pyridyl, 4-pyrimidinyl and 4-quinolyl, especially 4-pyridyl, are preferred. A preferred substituent for all R₁ moieties is C₁₋₄ alkyl, in particular methyl. More preferred as a substituted R₁ moiety is the 4-pyridyl derivative substituted at the 2-position with C₁₋₄ alkyl, especially 2-methyl-4-pyridyl. Also preferred is the 4-pyrimidinyl derivative substituted at the 2-position with C₁₋₄ alkyl or NR₁₀R₂₀.

20 Preferably, R₂ is hydrogen or C₁₋₁₀ alkyl, more preferably, hydrogen or methyl.

25 Preferably, the R₃ moieties is an (un)substituted aryl or heteroaryl moiety Q, also referred to as Q-(Y₁)_t. Preferably, when Q is an aryl, specifically phenyl, and when Q is a heteroaryl, preferred groups include pyrrole, pyridine, or pyrimidine. More preferred is Q as phenyl. All preferred moieties are independently substituted by (Y₁)_t, wherein t is an integer of 1 to 3. Preferably t is 1 or 2. More preferably, when R₃ is monosubstituted phenyl, the substituent is located at the 4-position.

30 Suitably the aryl or heteroaryl moiety of Q is substituted by up to three substituents Y₁ each of which is independently selected from C₁₋₅ alkyl, halo-substituted C₁₋₅ alkyl, halogen, -X_a-P(Z)-(X_bR₁₃)₂ or -(CR₁₀R₂₀)_nY₂ wherein Y₂ is -OR₈, -NO₂, -S(O)_mR₁₁, -SR₈, -S(O)_m'OR₈, -S(O)_mNR₈R₉, -NR₈R₉, -O(CR₁₀R₂₀)_nNR₈R₉, -C(O)R₈, -CO₂R₈, -CO₂(CR₁₀R₂₀)_n'CONR₈R₉, -ZC(O)R₈, -CN, -C(Z)NR₈R₉, -NR₁₀C(Z)R₈, -C(Z)NR₈OR₉, -NR₁₀C(Z)NR₈R₉, -NR₁₀S(O)_mR₁₁, -N(OR₂₁)C(Z)NR₈R₉, -N(OR₂₁)C(Z)R₈, -C(=NOR₂₁)R₈,

- NR₁₀C(=NR₁₅)SR₁₁, -NR₁₀C(=NR₁₅)NR₈R₉, -NR₁₀C(=CR₁₄R₂₄)SR₁₁,
- NR₁₀C(=CR₁₄R₂₄)NR₈R₉, -NR₁₀C(O)C(O)NR₈R₉, -NR₁₀C(O)C(O)OR₁₀,
- C(=NR₁₃)NR₈R₉, -C(=NOR₁₃)NR₈R₉, -C(=NR₁₃)ZR₁₁, -OC(Z)NR₈R₉,
- NR₁₀S(O)_mCF₃, -NR₁₀C(Z)OR₁₀, 5-(R₁₈)-1,2,4-oxadizaol-3-yl or 4-(R₁₂)-5-
- 5 (R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl; m' is 1 or 2; R₉ is hydrogen, C₁-10 alkyl, C₂-10 alkenyl, C₂-10 alkynyl, C₃-7 cycloalkyl, C₅-7 cycloalkenyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl or R₈ and R₉ may together with the nitrogen to which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected
- 10 from oxygen, sulfur or NR₁₂; R₁₄ and R₂₄ is each independently selected from hydrogen, alkyl, nitro or cyano; R₁₅ is hydrogen, cyano, C₁-4 alkyl, C₃-7 cycloalkyl or aryl; R₁₈ and R₁₉ is each independently selected from hydrogen, C₁-4 alkyl, substituted alkyl, optionally substituted aryl, arylalkyl or together with the carbon to which they are attached denote a double bonded oxygen or
- 15 sulfur, i.e., a C=O or C=S; and R₂₁ is hydrogen, a pharmaceutically acceptable cation, alkyl, cycloalkyl, aryl, arylalkyl, heteroaryl, heterocyclic, heteroarylalkyl, aroyl, alkoyl.

Preferred substituents Y₁ for use in R₃ include halogen, C₁-5 alkyl and -(CR₁₀R₂₀)_nY₂ wherein Y₂ is -OR₈, -NO₂, -S(O)_mR₁₁, -SR₈,

- 20 -S(O)_mNR₈R₉; -NR₈R₉, -O(CR₁₀R₂₀)_nNR₈R₉, -C(O)R₈, -CO₂R₈,
- CO₂(CR₁₀R₂₀)_nCONR₈R₉, -CN; -C(Z)NR₈R₉, -NR₁₀S(O)_mR₁₁,
- NR₁₀C(Z)R₈, -NR₁₀C(Z)NR₈R₉, -C(Z)NR₈OR₉, -N(OR₂₁)C(Z)NR₈R₉,
- NR₁₀C(=NR₁₅)NR₈R₉, -C(=NOR₁₃)NR₈R₉, 5-(R₁₈)-1,2,4-oxadizaol-3-yl and 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl.

- 25 Preferred substituents Y₁ for use in R₃ when the aryl or heteroaryl group Q is mono-substituted include -(CR₁₀R₂₀)_nY₂ wherein: n is 0, 1, 2 or 3, preferably 0 or 1; and Y₂ is -OR₈, especially where R₈ is hydrogen or C₁-10 alkyl; -NO₂; -S(O)_mR₁₁, especially where R₁₁ is C₁-10 alkyl; -SR₈, especially where R₈ is C₁-10 alkyl; -S(O)_mNR₈R₉, especially where R₈ and R₉ is each hydrogen or C₁-10 alkyl or R₈ and R₉ together with the nitrogen to which they are attached form a 5 to 7 membered ring which optionally includes another heteroatom selected from oxygen, sulfur or NR₁₂ and m is 2; n' is 1 to 10; -NR₈R₉, especially where R₈ and R₉ is each hydrogen, methyl or benzyl or R₈ and R₉ together with the nitrogen to which they are attached form a 5 to 7 membered ring which optionally includes another heteroatom selected from oxygen, sulfur or NR₁₂; -O(CR₁₀R₂₀)_nNR₈R₉, especially where R₈ and R₉ is each C₁-10 alkyl; -C(O)R₈, especially where R₈ is hydrogen or C₁-10 alkyl; -CO₂R₈, especially where R₈ is hydrogen or C₁-10 alkyl; -CO₂(CR₁₀R₂₀)_nCONR₈R₉, especially where R₈ and R₉ is

hydrogen or C₁₋₁₀ alkyl; -CN; -C(Z)NR₈R₉, especially where R₈ and R₉ is hydrogen or C₁₋₁₀ alkyl; -NR₁₀S(O)_mR₁₁, especially where R₁₀ is hydrogen or C₁₋₁₀ alkyl and R₁₁ is C₁₋₁₀ alkyl or a halosubstituted ; -NR₁₀C(Z)R₈, especially where R₈ is C₁₋₁₀ alkyl and R₁₀ is hydrogen and Z is oxygen;

5 -C(Z)NR₈OR₉, especially where R₈ and R₉ is each hydrogen and Z is oxygen; -NR₁₀C(Z)NR₈R₉, especially where R₈ and R₉ is each hydrogen or C₁₋₁₀ alkyl and Z is oxygen; -N(OR₂₁)C(Z)NR₈R₉, especially where R₈ especially where R₈, R₉ and R₂₁ is each hydrogen or C₁₋₁₀ alkyl and Z is oxygen; -C(=NOR₁₃)NR₈R₉, especially where R₈, R₉ and R₁₃ is each hydrogen;

10 -NR₁₀C(=NR₁₅)NR₈R₉, especially where R₈ and R₉ is hydrogen, C₁₋₁₀ alkyl or arylalkyl and R₁₅ is cyano; and 5-(R₁₈)-1,2,4-oxadizaol-3-yl and 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl, especially where R₁₂ is hydrogen and R₁₈ and R₁₉ is each hydrogen or C₁₋₁₀ alkyl or together are oxo.

15 Preferred substituents for use in R₃ when the aryl or heteroaryl group Q is disubstituted include those hereinbefore listed for use when Q is mono-substituted and, as further substituent(s), halogen and C₁₋₁₀ alkyl. When R₃ is phenyl substituted with two or three substituents, the alkyl moieties preferably have from one to three carbons, more preferably one. Preferred 20 ring positions for two substituents are the 3- and 4-positions and, for three substituents, the 3-, 4- and 5- positions. The substituent at the 3- and 5- positions is preferably C₁₋₂ alkyl, such as methyl, or halogen, such as bromo, fluoro or chloro, while the substituent at the 4-position is preferably hydroxyl.

More preferably, for R₃ substituents wherein Y₁ is (CR₁₀R₂₀)_nY₂, n is 0 or 1 and Y₂ is -OH, -S(O)_mR₁₁, especially where R₁₁ is C₁₋₁₀ alkyl; -SR₈, especially where R₈ is C₁₋₁₀ alkyl; -NR₈R₉, especially where R₈ and R₉ is hydrogen, alkyl, aryl alkyl, or aryl or R₈ and R₉ together with the nitrogen to which they are attached form a pyrrolidinyl, piperidinyl or morpholinyl ring, more prefereably the R₈ and R₉ terms in the NR₈R₉ 30 moiety are hydrogen, methyl or benzyl; -CO₂R₈, especially where R₈ is hydrogen or C₁₋₁₀ alkyl; -S(O)_mNR₈R₉, especially where R₈ and R₉ is each hydrogen or C₁₋₁₀ alkyl; -NR₁₀S(O)_mR₁₁, especially where R₁₀ is hydrogen and R₁₁ is C₁₋₁₀ alkyl or 5-(R₁₈)-1,2,4-oxadizaol-3-yl and 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl, especially where R₁₂ is hydrogen and R₁₈ 35 and R₁₉ is hydrogen or C₁₋₁₀ alkyl or together are oxo.

Most preferably, Y₁ is methylthio, ethylthio, methylsulfinyl, ethylsulfinyl, methylsulfonyl, N,N-dimethylaminomethyl, N-benzyl-N-

methylaminomethyl, N-morpholinomethyl, methanesulfonamido, sulphonamidomethyl, 5-methyl-4,5-dihydro-1,2,4-oxadiazol-3-yl or 5,5-dimethyl-4,5-dihydro-1,2,4-oxadiazol-3-yl.

In all instances herein where there is an alkenyl or alkynyl moiety as 5 a substituent group, such as in R₅, R₈, R₉, or R₁₁ the unsaturated linkage, i.e., the vinylene or acetylene linkage is preferably not directly attached to the nitrogen, oxygen or sulfur moieties, for instance in Y₂ as C(Z)NR₈OR₉, NR₁₀C(Z)NR₈R₉, or OR₈. As used herein, "optionally substituted" unless specified refers to such groups as halogen, hydroxyl, alkoxy, S(O)_m alkyl, 10 amino, mono & di-substituted amino, such as a NR₇R₁₇ group, alkyl or cycloalkyl, i.e. such as in optionally substituted aryl or optionally substituted arylalkyl.

When R₃ includes a X_a-P(Z)(X_bR₁₃)₂ group linked either directly to the imidazole ring or indirectly *via* an aryl or heteroaryl group, X_a is 15 suitably oxygen or C₁₋₄ alkylene, optionally interrupted by oxygen, for instance -CH₂OCH₂- and Z and X_b is each oxygen, such that the preferred groups include -OP(O)(OR₁₃)₂ and -CH₂OCH₂-P(O)(OR₁₃)₂.

Preferred substitutions for R₄ when this is a 4-phenyl, 4-naphth-1-yl or 5-naphth-2-yl moiety are one or two substituents each independently 20 selected from halogen, -SR₅, -SOR₅, -OR₃₆, or -(CR₁₀R₂₀)_mNR₁₀R₂₀, and for other positions of substitution on these rings preferred substitution is halogen, -S(O)_mR₈, -OR₈, -(CR₁₀R₂₀)_mNR₁₆R₂₆, -NR₁₀C(Z)R₈ and -NR₁₀S(O)_mR₁₁. More preferred substituents for the 4-position in phenyl and naphth-1-yl and on the 5-position in naphth-2-yl include halogen, 25 especially fluoro and chloro, and -SR₅ and -SOR₅ wherein R₅ is preferably a C₁₋₂ alkyl, more preferably methyl; of which fluoro is especially preferred. Preferred substituents for the 3-position in phenyl and naphth-1-yl include: halogen, especially chloro; -OR₈, especially C₁₋₄ alkoxy; amino; -NR₁₀C(Z)R₈, especially -NHCO(C₁₋₁₀ alkyl); and -NR₁₀S(O)_mR₁₁, 30 especially -NHSO₂(C₁₋₁₀ alkyl). Preferably, the R₄ moiety is an unsubstituted or substituted phenyl moiety. More preferably, R₄ is phenyl or phenyl substituted at the 4-position with fluoro and/or substituted at the 3-position with fluoro, chloro, C₁₋₄ alkoxy, methanesulfonamido or acetamido.

In a preferred subgenus of compounds of formula (I), R₁ is 4-pyridyl, 35 2-alkyl-4-pyridyl or 4-quinolyl; R₂ is hydrogen or methyl; R₃ is phenyl or phenyl substituted, preferably at the 4-position, with a substituent selected from -(CR₁₀R₂₀)_nY₂ wherein Y₂ is wherein n is 0, 1, 2 or 3 and Y₂ is -OR₈, -NO₂, -S(O)_mR₁₁, -SR₈, -S(O)_mNR₈R₉, -NR₈R₉, -O(CR₁₀R₂₀)_nNR₈R₉,

-C(O)R₈, -CO₂R₈, -CO₂(CR₁₀R₂₀)_nCONR₈R₉, -CN, -C(Z)NR₈R₉, -C(Z)NR₈OR₉, -NR₁₀S(O)_mR₁₁, -NR₁₀C(Z)R₈, -NR₁₀C(Z)NR₈R₉, -C(=NOR₁₃)NR₈R₉, -NR₁₀C(=CR₁₄R₂₄)NR₈R₉, 5-(R₁₈)-1,2,4-oxadizaol-3-yl, 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl, a 3,5-dimethyl or 5 dibromo-4-hydroxyl grouping; and R₄ is phenyl or phenyl substituted at the 4-position with fluoro and/or substituted at the 3-position with fluoro, chloro, C₁₋₄ alkoxy, methanesulfonamido or acetamido.

In a more preferred subgenus, R₁ is 4-pyridyl, 2-methyl-4-pyridyl or 4-quinolyl; R₂ is hydrogen or methyl; R₃ is phenyl substituted at the 4-position with C₁₋₁₀ alkylthio, C₁₋₁₀ alkylsulfinyl, C₁₋₁₀ alkylsulfonyl, N,N-di(C₁₋₁₀ alkyl)amino C₁₋₂ alkyl, N-aralkyl-N-C₁₋₁₀ alkylamino C₁₋₂ alkyl, N-morpholino C₁₋₂ alkyl, C₁₋₁₀ alkylsulfonamido, sulphonamido C₁₋₂ alkyl, 5-C₁₋₁₀ alkyl-4,5-dihydro-1,2,4-oxadiazol-3-yl or 5,5-di(C₁₋₁₀ alkyl)-4,5-dihydro-1,2,4-oxadiazol-3-yl; and R₄ is phenyl or phenyl substituted at the 4-position with fluoro and/or substituted at the 3-position with fluoro, chloro, C₁₋₄ alkoxy, methanesulfonamido or acetamido.

Suitable pharmaceutically acceptable salts are well known to those skilled in the art and include basic salts of inorganic and organic acids, such as hydrochloric acid, hydrobromic acid, sulphuric acid, phosphoric acid, 20 methane sulphonic acid, ethane sulphonic acid, acetic acid, malic acid, tartaric acid, citric acid, lactic acid, oxalic acid, succinic acid, fumaric acid, maleic acid, benzoic acid, salicylic acid, phenylacetic acid and mandelic acid. In addition, pharmaceutically acceptable salts of compounds of formula (I) may also be formed with a pharmaceutically acceptable cation, for instance, 25 if a substituent Y₁ in R₃ comprises a carboxy group. Suitable pharmaceutically acceptable cations are well known to those skilled in the art and include alkaline, alkaline earth, ammonium and quarternary ammonium cations.

The following terms, as used herein, refer to:

30 • "halo" - all halogens, that is chloro, fluoro, bromo and iodo;

• "C₁₋₁₀alkyl" or "alkyl" - both straight and branched chain radicals of 1 to 10 carbon atoms, unless the chain length is otherwise limited, including, but not limited to, methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *sec*-butyl, *iso*-butyl, *tert*-butyl, and the like;

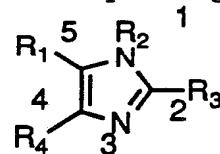
35 • "aryl" - phenyl and naphthyl;

• "heteroaryl" (on its own or in any combination, such as "heteroaryloxy") - a 5-10 membered aromatic ring system in which one or more rings contain one or more heteroatoms selected from the group

consisting of N, O or S, such as, but not limited, to pyrrole, quinoline, isoquinoline, pyridine, pyrimidine, oxazole, thiazole, thiadiazole, triazole, imidazole, or benzimidazole;

- "heterocyclic" (on its own or in any combination, such as
 - 5 "heterocyclalkyl" - a saturated or wholly or partially unsaturated 4-10 membered ring system in which one or more rings contain one or more heteroatoms selected from the group consisting of N, O, or S; such as, but not limited to, pyrrolidine, piperidine, piperazine, morpholine, imidazolidine or pyrazolidine;
 - 10 • "aryloyl" - a C(O)Ar, wherein Ar is as phenyl, napthyl, or aryl alkyl derivatives, such as benzyl and the like;
 - "alkoyl" - a C(O)C₁₋₁₀alkyl wherein the alkyl is as defined above;
 - "sulfinyl" - the oxide (SO) of the corresponding sulfide whilst the term "thio" refers to the sulfide.
- 15 The compounds of the present invention may contain one or more asymmetric carbon atoms and may exist in racemic and optically active forms. All of these compounds are included within the scope of the present invention.

For the purposes herein of nomenclature, the compounds of formula (I) are named by their position corresponding to:



Especially preferred compounds of formula (I) include:

- 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole;
- 25 4-(4-Fluorophenyl)-2-(4-ethylthiophenyl)-5-(4-pyridyl)imidazole;
- 4-(4-Fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)imidazole;
- 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)imidazole;
- 4-(4-Fluorophenyl)-2-(4-ethylsulfinylphenyl)-5-(4-pyridyl)imidazole;
- 4-(3-Chlorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)imidazole;
- 30 2-[4-(N-Methyl-N-benzyl)aminomethylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)imidazole;
- 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)imidazole;
- 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylsulfinylphenyl)-imidazole;
- 35 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-quinolyl)imidazole;

2-[4-(N-Morpholino)methylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-imidazole; and pharmaceutically acceptable salts thereof.

5 Other preferred compounds of formula (I) include:

2-[(4-N,N-Dimethyl)aminomethylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-imidazole;

2-(4-Methanesulfonamidophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)imidazole;

4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-quinolyl)imidazole;

10 4-(3-Chlorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole;

4-(3-Methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole;

4-(3-Methoxyphenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)imidazole;

4-(3-Methanesulfonamidophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-imidazole;

15 4-(3-Methanesulfonamidophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-imidazole;

3-[4-(4-Fluorophenyl)-5-(4-pyridyl)imidazol-2-yl]phenyl-5,5-dimethyl-4,5-dihydro-1,2,4-oxadiazole; or

3-[4-(4-Fluorophenyl)-5-(4-pyridyl)imidazol-2-yl]phenyl-5-methyl-4,5-dihydro-1,2,4-oxadiazole; and

20 pharmaceutically acceptable salts thereof.

Compounds of formula (I) are imidazole derivatives which may be readily prepared using procedures well-known to those skilled in the art, 25 and described in, for instance, Comprehensive Heterocyclic Chemistry, ed Katritzky and Rees, Pergamon Press, 1984, 5, 457-497, from starting materials which are either commercially available or can be prepared from such by analogy with well-known processes. A key step in many such syntheses is the formation of the central imidazole nucleus, to give 30 compounds of formula (I). Suitable procedures are described in *inter alia* US patent nos. 3,707,475 and 3,940,486 which are herein incorporated by reference in their entirety. These patents describe the synthesis of α -diketones and α -hydroxyketones (benzoins) and their subsequent use in preparing imidazoles and N-hydroxyl imidazoles. Thereafter, further 35 compounds of formula (I) may be obtained by manipulating substituents in any of the groups R₁, R₂, R₃ and R₄ using conventional functional group interconversion procedures.

In particular, in a first process, compounds of formula (I) may be prepared by condensing an α -diketone of formula (II):



wherein R_1 and R_4 are as hereinbefore defined, or an equivalent thereof, with an aldehyde of the formula (III):



5 wherein R_3 is as hereinbefore defined, or an equivalent thereof, and, if necessary, with ammonia or a source thereof, under imidazole-ring forming conditions.

Suitable equivalents of the α -diketone are well known to those skilled in the art and include the corresponding α -keto-oxime and α -dioxime.

10 Suitable equivalents of the aldehyde of formula (III) are well known in the art and include the corresponding oxime and acetal.

Ammonia, or a source thereof, is preferably used in excess, with at least a dimolar amount being used in the case of the α -diketone and at least an equimolar amount in the case of the α -keto-oxime.

15 Suitable sources of ammonia include ammonium salts of organic carboxylic acids, such as an ammonium C₁-6 alkanoate, for instance ammonium acetate and ammonium formate, preferably ammonium acetate, and carboxylic amides, in particular of formic acid, such as formamide. An ammonium salt is generally used in large excess and in the presence of an acid, such as a C₁-6 carboxylic acid which acid may also be used as a solvent for the reaction. If formamide is used, this may be used in excess, as the reaction solvent. An alternative solvent such as ethanol or dimethyl sulphoxide (Lantos *et al*, J Het Chem, 19, 1375, 1982) may be used. An additional solvent may also be employed, for instance, dimethyl formamide

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may be used with formamide. The reaction is generally carried out at elevated temperatures, for instance under reflux conditions, and if desired, in a sealed vessel optionally under pressure and/or an inert gas atmosphere, for instance nitrogen.

A further suitable source of ammonia is hydroxylamine, in which case

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35

the initially formed imidazole is an N-hydroxy-N-oxide imidazole. This may then be reduced to the corresponding N-hydroxy imidazole by treating with a suitable reducing agent such as sodium borohydride, in an appropriate solvent such as methanol, following the method of Akange and Allan, Chem and Ind, 5 Jan 1975, 38. The N-hydroxy imidazole may in turn be converted to an imidazole of formula (I) in which R_2 is hydrogen by treatment with a conventional deoxygenating agent such as phosphorus trichloride or a trialkylphosphite such as trimethyl- or triethyl-phosphite. N-hydroxy-N-oxide imidazoles may be readily obtained by treating an α -diketone of

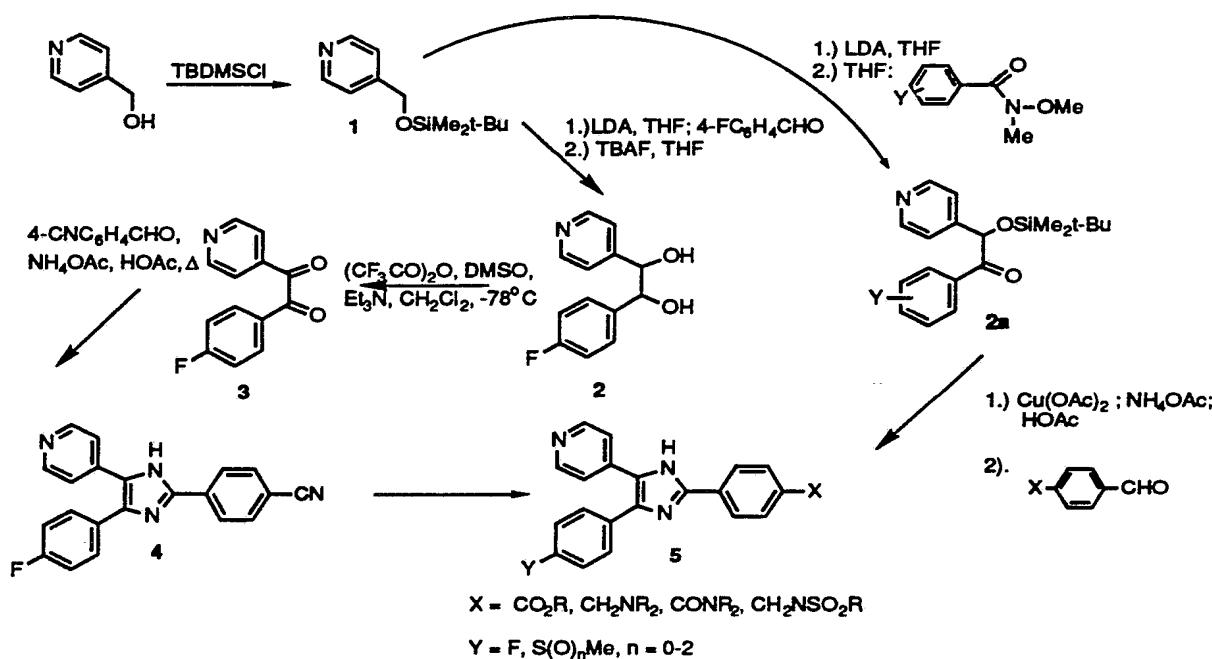
formula (II) with an aldehyde of formula (II) with about two equivalents of hydroxylamine or the corresponding aldoxime and about one equivalent of hydroxylamine, under proton catalysis. Alternatively, the N-oxide may be obtained by the acid catalysed condensation of the corresponding α -dioxime or α -keto-oxime with an aldoxime of the aldehyde of formula (III).

5 When the compound of formula (II) is an α -keto-oxime derivative, it will be appreciated that the product initially obtained will be a compound of formula (I) in which R_2 is hydroxyl which may be converted into a compound of formula (I) in which R_2 is hydrogen as described above.

10 It will be appreciated by those skilled in the art that in some instances, it will not be necessary to provide a separate source of ammonia as the α -diketone or aldehyde equivalent may already contain such a source. Examples of this include α -dioxime or α -keto-oxime and aldoxime.

15 The compounds of formula (II) may be obtained by applying well-known synthetic procedures, some of which are illustrated in schemes I and II. Although these illustrate syntheses in which R_4 is either 4-pyridyl or 4-quinolinyl, they may be equally applied to any of the other heteroaryl rings within the definition of R_4 by appropriate choice of starting material.

20 In Scheme I, the anion prepared from 1, by treatment with a strong base such as lithium di-*iso*-propylamide, is condensed with a substituted benz-aldehyde, to give, after removal of the protecting group, the diol 2. This may then be converted to the α -diketone 3 by a Swern oxidation of which any number of potentially useful variations are known and may be used. The α -diketone 3 is then cyclised to an imidazole 4, a compound of 25 formula (I), by heating 3 with a substituted benzaldehyde in a mixture of ammonium acetate, as the source of ammonia, and an appropriate solvent, for example acetic acid or DMSO. The imidazole 4 may then be transformed into other imidazoles 5 by appropriate functional group 30 interconversion procedures. Scheme I also illustrates the preparation of a protected α -hydroxyketone 2a, by condensing the anion of 1 with an appropriately activated carbonyl derivative of a substituted benzamide, such as the N-methoxy-N-methylamide, to yield a protected α -hydroxyketone. This adduct 2a may then be directly converted to the imidazole 5, using a combination of a copper (II) salt, such as copper (II) acetate, as an oxidising agent and ammonium acetate as a source of 35 ammonia. The α -hydroxyketone 2a may also be deprotected and then oxidised to give an α -diketone 3, for instance using Swern oxidation.

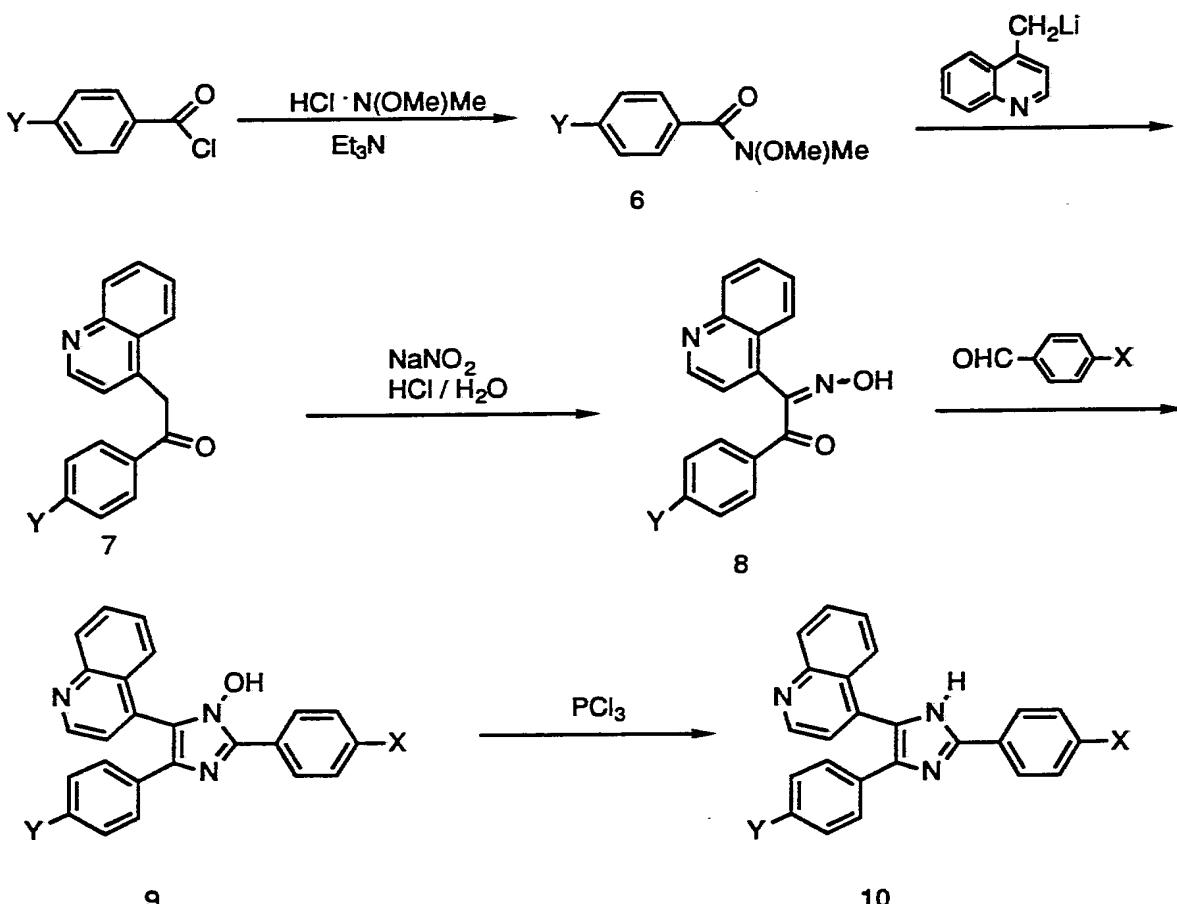


Scheme I

5 Scheme II illustrates the use of an α -keto-oxime for preparing a compound of formula (I). A heterocyclic ketone 7 is prepared by adding the anion of 4-methyl-quinoline (prepared by treatment thereof with an alkyl lithium, such as *n*-butyl lithium) to an **N**-alkyl-**O**-alkoxybenzamide. Alternatively, the anion may be condensed with a benzaldehyde, to give an alcohol which is then oxidised to the ketone 7. The α -keto-oxime 8 is then prepared from 7 using standard conditions, such as reaction with sodium nitrite, and this may then be reacted with a benzaldehyde to afford an **N**-hydroxyimidazole 9, a compound of formula (I) in which R_2 is hydroxy. This may be converted to 10, a further compound of formula (I) in which R_2 is

10 hydrogen, by treating it with a deoxygenating agent such as phosphorus trichloride or a trialkyl phosphite, such as trimethyl or triethylphosphite. For compounds of formula (I) wherein R_3 is $-(\text{CR}_{10}\text{R}_{20})_n\text{-P}(\text{Z})-(\text{X}_b\text{R}_{13})_2$, the reagent $\text{OHC}-(\text{CR}_{10}\text{R}_{20})_n\text{-P}(\text{Z})-(\text{X}_b\text{R}_{13})_2$ may be used instead of $\text{OHC-C}_6\text{H}_4\text{-X}$ to make the appropriately substituted compound 9.

15



Scheme II

5 In a further process, a compound of formula (I) may be obtained by
treating an α -hydroxyketone compound of formula (IIA):



10 wherein one of R' and R'' is R₁ and the other is R₄, a suitably protected
derivative thereof or the α -hydroxy-oxime or α -haloketone derivative
thereof, with an oxidising agent capable of converting said compound into
the corresponding α -diketone, in the presence of an aldehyde of formula (III)
or an equivalent thereof, and a source of ammonia. Suitable oxidising
agents include, for example, an oxidising heavy metal salt, preferably an
organic copper (II) salt, such as copper (II) acetate or copper (II) citrate. The
15 reaction may be effected in a solvent such as acetic acid, under reflux
conditions. Alternatively, a lower alkanol solvent, such as methanol or
ethanol, may be used, preferably at a temperature in the region of from 30
to 100°C (see *The Chemistry of Heterocyclic Compounds, Imidazole and its*

derivatives, part I, ed. Weissberger, Interscience Publishers, Inc., New York, 1953, 38). This approach is also illustrated in Scheme I.

In a further process, a compound of formula (I) may be obtained by treating an amidine of formula (IV):

5



wherein R_2 and R_3 are as hereinbefore defined, or a salt thereof, with a reactive ester of an α -hydroxyketone of formula (IIA) or the corresponding α -haloketone, in an inert solvent such as a halogenated hydrocarbon solvent, for example chloroform, at a moderately elevated temperature and, 10 if necessary, in the presence of a suitable condensation agent such as a base. Suitable reactive esters include esters of strong organic acids such as a lower alkane sulphonic or aryl sulphonic acid, for instance, methane or *p*-toluene sulphonic acid. The amidine of formula (IV) is preferably used as the salt, suitably the hydrochloride salt, which may then be converted into 15 the free amidine *in situ*, by employing a two phase system in which the reactive ester is in an inert organic solvent such as chloroform, and the salt is in an aqueous phase to which a solution of an aqueous base is slowly added, in dimolar amount, with vigorous stirring. Suitable amidines of formula (IV) may be obtained by standard methods, see for instance, 20 Garigipati R, Tetrahedron Letters, 190, 31, 1989.

In a further process, a compound of formula (I) may be obtained by treating an iminoether of formula (V):



wherein R_3 is as hereinbefore defined and R is C_{1-10} alkyl, aryl or aryl C_{1-4} 25 alkyl, with an α -aminoketone of the formula (VI):



wherein one of R' and R'' is R_1 and the other is R_4 in a suitable solvent.

In a further process, N-substituted compounds of formula (I) may be prepared by treating the anion of an amide of formula (VII):

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wherein R_1 and R_3 are as hereinbefore defined and R_2 is as hereinbefore defined other than hydrogen, with:

(a) a nitrile of the formula (VIII):



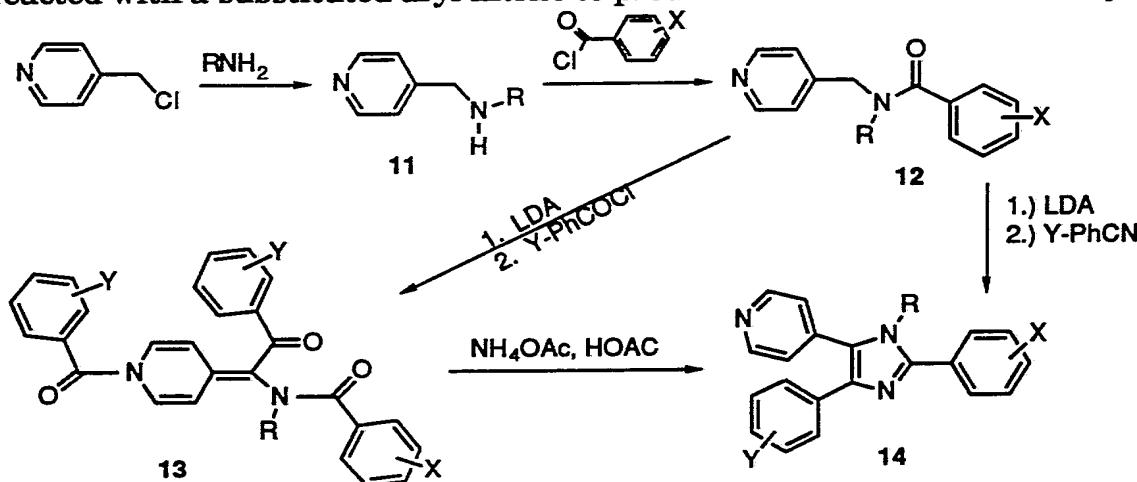
35 wherein R_4 is as hereinbefore defined, or

(b) an excess of an acyl halide, for instance an acyl chloride, of the formula (IX):



wherein R_4 is as hereinbefore defined and Hal is halogen, or a corresponding anhydride, to give a *bis*-acylated intermediate which is then treated with a source of ammonia, such as ammonium acetate.

This approach permits the regiospecific preparation of compound of formula (I) substituted at the 1-position, as illustrated in Scheme III. A primary amine RNH_2 is treated with 4-chloromethylpyridine to give 11 which is then converted to the amide 12 by standard techniques. Deprotonation of 12 with a strong amide base, such as lithium di-*iso*-propyl amide or sodium *bis*-(trimethylsilyl)amide, followed by addition of an excess of an aroyl chloride yields the *bis*-acylated compound 13 which is then closed to an imidazole compound of formula (I), 14, by heating in acetic acid containing ammonium acetate. Alternatively, the anion of 12 may be reacted with a substituted aryl nitrile to produce the imidazole 14 directly.



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Scheme III

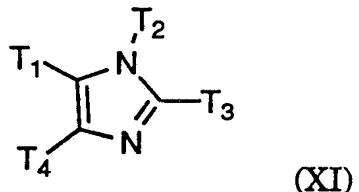
In a further process, compounds of formula (I) may be prepared by treating a compound of formula (X):



wherein R' , R'' and R_3 are as hereinbefore defined and X_c is O or NH, with a source of ammonia, as hereinbefore described, under imidazole ring forming conditions or cyclising the corresponding Schiff's base, formed by treating the compound of formula (X) in which X_c is NH with an amine R_2NH_2 , for instance thermally or with the aid of a cyclising agent such as phosphorus oxychloride or phosphorus pentachloride (see Engel and Steglich, Liebigs Ann Chem, 1978, 1916 and Strzybny *et al.*, J Org Chem, 1963, 28, 3381). Compounds of formula (X) may be obtained, for instance, by acylating the corresponding α -keto-oxime (X_c is NH) or α -hydroxyketone (X_c is O) with an

acyl halide of the formula R_3COHal wherein R_3 is as hereinbefore defined, or the corresponding anhydride, under standard acylating conditions.

In a further process, compounds of formula (I) may be prepared by coupling a suitable derivative of a compound of formula (XI):



wherein: T_2 is a nitrogen protecting group or R_2 , other than hydrogen; and T_1 is hydrogen, T_3 is Q and T_4 is R_4 ; T_1 is R_1 , T_3 is hydrogen and T_4 is R_4 ; or T_1 is R_1 , T_3 is Q and T_4 is hydrogen, in which R_1 , R_2 , R_3 , R_4 and Q are as hereinbefore defined; with: (i) when T_1 is hydrogen, a suitable derivative of the heteroaryl ring R_1H , under ring coupling conditions, to effect coupling of the heteroaryl ring R_1 to the imidazole nucleus at position 5; (ii) when T_3 is hydrogen, a suitable derivative of the aryl or heteroaryl ring QH , under ring coupling conditions, to effect coupling of the ring Q to the imidazole nucleus at position 2; or (iii) when T_4 is hydrogen, a suitable derivative of the aryl ring R_4H , under ring coupling conditions, to effect coupling of the aryl ring R_4 to the imidazole nucleus at position 4.

Such aryl/heteroaryl coupling reactions are well known to those skilled in the art. In general, an organometallic synthetic equivalent of an anion of one component is coupled with a reactive derivative of the second component, in the presence of a suitable catalyst. The anion equivalent may be formed from either the imidazole of formula (XI), in which case the aryl/heteroaryl compound provides the reactive derivative, or the aryl/heteroaryl compound in which case the imidazole provides the reactive derivative. Accordingly, suitable derivatives of the compound of formula (XI) or the aryl/heteroaryl rings include organometallic derivatives such as organomagnesium, organozinc, organostannane and boronic acid derivatives and suitable reactive derivatives include the the bromo, iodo, fluorosulfonate and trifluoromethanesulphonate derivatives. Suitable procedures are described in WO 91/19497, the disclosure of which is herewith incorporated.

30 Suitable organomagnesium and organozinc derivatives of a compound of formula (XI) may be reacted with a halogen, fluorosulfonate or triflate derivative of the heteroaryl or aryl ring, in the presence of a ring coupling catalyst, such as a palladium (0) or palladium (II) catalyst, following the procedure of Kumada *et al.*, *Tetrahedron Letters*, **22**, 5319 (1981). Suitable 35 such catalysts include *tetrakis*-(triphenylphosphine)palladium and

PdCl₂[1,4-*bis*-(diphenylphosphino)-butane], optionally in the presence of lithium chloride and a base, such as triethylamine. In addition, a nickel (II) catalyst, such as Ni(II)Cl₂(1,2-biphenylphosphino)ethane, may also be used for coupling an aryl ring, following the procedure of Pridgen, J Org Chem, 1982, 47, 4319. Suitable reaction solvents include hexamethylphosphoramide. When the heteroaryl ring is 4-pyridyl, suitable derivatives include 4-bromo- and 4-iodo-pyridine and the fluorosulfonate and triflate esters of 4-hydroxy pyridine. Similarly, suitable derivatives for when the aryl ring is phenyl include the bromo, fluorosulfonate, triflate and, preferably, the iodo-derivatives. Suitable organomagnesium and organozinc derivatives may be obtained by treating a compound of formula (XI) or the bromo derivative thereof with an alkyl lithium compound to yield the corresponding lithium reagent by deprotonation or transmetallation, respectively. This lithium intermediate may then be treated with an excess of a magnesium halide or zinc halide to yield the corresponding organometallic reagent.

A trialkyltin derivative of the compound of formula (XI) may be treated with a bromide, fluorosulfonate, triflate, or, preferably, iodide derivative of an aryl or heteroaryl ring compound, in an inert solvent such as tetrahydrofuran, preferably containing 10% hexamethylphosphoramide, in the presence of a suitable coupling catalyst, such as a palladium (0) catalyst, for instance *tetrakis*-(triphenylphosphine)palladium, by the method described in by Stille, J Amer Chem Soc, 1987, 109, 5478, US Patents 4,719,218 and 5,002,942, or by using a palladium (II) catalyst in the presence of lithium chloride optionally with an added base such as triethylamine, in an inert solvent such as dimethyl formamide. Trialkyltin derivatives may be conveniently obtained by metallation of the corresponding compound of formula (XI) with a lithiating agent, such as *s*-butyl-lithium or *n*-butyllithium, in an ethereal solvent, such as tetrahydrofuran, or treatment of the bromo derivative of the corresponding compound of formula (XI) with an alkyl lithium, followed, in each case, by treatment with a trialkyltin halide. Alternatively, the bromo- derivative of a compound of formula (XI) may be treated with a suitable heteroaryl or aryl trialkyl tin compound in the presence of a catalyst such as *tetrakis*-(triphenylphosphine)-palladium, under conditions similar to those described above.

Boronic acid derivatives are also useful. Hence, a suitable derivative of a compound of formula (XI), such as the bromo, iodo, triflate or fluorosulphonate derivative, may be reacted with a heteroaryl- or aryl-boronic acid, in the presence of a palladium catalyst such as *tetrakis*-(triphenylphosphine)-palladium or PdCl₂[1,4-*bis*-(diphenylphosphino)-

butane] in the presence of a base such as sodium bicarbonate, under reflux conditions, in a solvent such as dimethoxyethane (see Fischer and Haviniga, Rec. Trav. Chim. Pays Bas, **84**, 439, 1965, Snieckus, V., Tetrahedron Lett., **29**, 2135, 1988 and Terashimia, M., Chem. Pharm. Bull., **11**, 4755, 1985).

5 Non-aqueous conditions, for instance, a solvent such as DMF, at a temperature of about 100°C, in the presence of a Pd(II) catalyst may also be employed (see Thompson W J *et al*, J Org Chem, **49**, 5237, 1984). Suitable boronic acid derivatives may be prepared by treating the magnesium or lithium derivative with a trialkylborate ester, such as triethyl, tri-*iso*-propyl or tributylborate, according to standard procedures.

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In such coupling reactions, it will be readily appreciated that due regard must be exercised with respect to functional groups present in the compounds of formula (XI). Thus, in general, amino and sulfur substituents should be non-oxidised or protected and the N-1 nitrogen of a compound of formula (XI) be protected, if an NH compound is finally required. Nitro, bromo, iodo and hydroxyl groups should preferably be avoided in compounds of formula (XI) in which T₁ is hydrogen.

Compounds of formula (XI) are imidazoles and may be obtained by any of the procedures herein before described for preparing compounds of formula (I). In particular, an α -halo-ketone R₄COCH₂Hal (for compounds of formula (XI) in which T₁ is hydrogen) or R₁COCH₂Hal (for compounds of formula (XI) in which T₄ is hydrogen) may be reacted with an amidine of formula (IV) or a salt thereof, in an inert solvent such as a halogenated hydrocarbon solvent, for instance chloroform, at a moderately elevated temperature, and, if necessary, in the presence of a suitable condensation agent such as a base. The preparation of suitable α -halo-ketones is described in WO 91/19497. For a compound of formula (XI) in which T₃ is hydrogen, an α -diketone of formula (II) may be condensed with a formaldehyde or an equivalent thereof, in the presence of a source of ammonia. Suitable bromo derivatives of the compound of formula (XI) may be obtained by brominating the corresponding compound of formula (XI) under standard brominating conditions, for instance bromine in a solvent such as dichloromethane or THF.

Compounds of formula (I) may also be prepared by a process which comprises reacting a compound of formula (XI), wherein T₁ is hydrogen, with an N-acyl heteroaryl salt, according to the method disclosed in US patents 4,803,279, 4,719,218 and 5,002,942, to give an intermediate in which the heteroaryl ring is attached to the imidazole nucleus and is present as a 1,4-dihydro derivative thereof, which intermediate may then be

subjected to oxidative-deacylation conditions. The heteroaryl salt, for instance a pyridinium salt, may be either preformed or, more preferably, prepared *in situ* by adding a substituted carbonyl halide (such as an acyl halide, an aroyl halide, an arylalkyl haloformate ester, or, preferably, an alkyl haloformate ester, such as acetyl bromide, benzoylchloride, benzyl chloroformate, or, preferably, ethyl chloroformate) to a solution of the compound of formula (XI) in the heteroaryl compound R₁H or in an inert solvent such as methylene chloride to which the heteroaryl compound has been added. Suitable deacylating and oxidising conditions are described in U.S. Patent Nos. 4,803,279, 4,719,218 and 5,002,942, which references are hereby incorporated in their entirety. Suitable oxidising systems include sulfur in an inert solvent or solvent mixture, such as decalin, decalin and diglyme, *p*-cymene, xylene or mesitylene, under reflux conditions, or, preferably, potassium *t*-butoxide in *t*-butanol with dry air or oxygen.

Once the imidazole nucleus has been established, further compounds of formula (I) which may be prepared by applying standard techniques for functional group interconversion, for instance: -C(O)NR₈R₉ from -CO₂CH₃ by heating with or without catalytic metal cyanide, e.g. NaCN, and HNR₈R₉ in CH₃OH; -OC(O)R₈ from -OH with e.g., ClC(O)R₈ in pyridine; -NR₁₀-C(S)NR₈R₉ from -NHR₁₀ with an alkylisothiocyanate or thiocyanic acid; NR₆C(O)OR₆ from -NHR₆ with the alkyl chloroformate; -NR₁₀C(O)NR₈R₉ from -NHR₁₀ by treatment with an isocyanate, e.g. HN=C=O or R₁₀N=C=O; -NR₁₀-C(O)R₈ from -NHR₁₀ by treatment with Cl-C(O)R₈ in pyridine; -C(=NR₁₀)NR₈R₉ from -C(NR₈R₉)SR₈ with H₃NR₈⁺OAc⁻ by heating in alcohol; -C(NR₈R₉)SR₈ from -C(S)NR₈R₉ with R₆-I in an inert solvent, e.g. acetone; -C(S)NR₈R₉ (where R₈ or R₉ is not hydrogen) from -C(S)NH₂ with HNR₈R₉; -C(=NCN)-NR₈R₉ from -C(=NR₈R₉)-SR₈ with NH₂CN by heating in anhydrous alcohol, alternatively from -C(=NH)-NR₈R₉ by treatment with BrCN and NaOEt in EtOH; -NR₁₀-C(=NCN)SR₈ from -NHR₁₀ by treatment with (R₈S)₂C=NCN; -NR₁₀SO₂R₈ from -NHR₁₀ by treatment with ClSO₂R₈ by heating in pyridine; -NR₁₀C(S)R₈ from -NR₁₀C(O)R₈ by treatment with Lawesson's reagent [2,4-bis(4-methoxyphenyl)-1,3,2,4-dithiadiphosphetane-2,4-disulfide]; -NR₁₀SO₂CF₃ from -NHR₆ with triflic anhydride and base; -NR₁₀C(O)-C(O)-OR₈ from -NHR₁₀ with, e.g. methyloxalyl chloride and a base such as triethylamine; -NR₁₀C(O)-C(O)-NR₈R₉ from -NR₁₀C(O)-C(O)-OR₈ with HNR₈R₉; and 1-(NR₁₀)-2-imidazolyl from -C(=NH)NHR₁₀ by heating with 2-chloroacetaldehyde in chloroform (wherein R₆, R₈, R₉ and R₁₀ are as hereinbefore defined).

Compounds of formula (I) in which R₂ is hydrogen may be readily converted into further compounds of formula (I) in which R₂ is other than hydrogen, for instance alkyl, by conventional procedures such as alkylation or acylation followed by reduction. Such methods are in general relatively inefficient as they lack regiospecificity and the desired N-1 product has to be separated from the mixture of N-1 and N-3 products, for instance by chromatography or fractional crystallisation.

Suitable protecting groups for use with hydroxyl groups and the imidazole nitrogen are well known in the art and described in many references, for instance, Protecting Groups in Organic Synthesis, Greene T W, Wiley-Interscience, New York, 1981. Suitable examples of hydroxyl protecting groups include silyl ethers, such as t-butyldimethyl or t-butyldiphenyl, and alkyl ethers, such as methyl connected by an alkyl chain of variable link, (CR₁₀R₂₀)_n. Suitable examples of imidazole nitrogen protecting groups include tetrahydropyranyl.

Pharmaceutically acid addition salts of compounds of formula (I) may be obtained in known manner, for example by treatment thereof with an appropriate amount of acid in the presence of a suitable solvent.

20 METHODS OF TREATMENT

The compounds of Formula (I) or a pharmaceutically acceptable salt thereof can be used in the manufacture of a medicament for the prophylactic or therapeutic treatment of any disease state in a human, or other mammal, which is exacerbated or caused by excessive or unregulated cytokine production by such mammal's cell, such as but not limited to monocytes and/or macrophages.

Compounds of formula (I) are capable of inhibiting proinflammatory cytokines, such as IL-1, IL-6, IL-8 and TNF and are therefore of use in therapy. IL-1, IL-8 and TNF affect a wide variety of cells and tissues and these cytokines, as well as other leukocyte-derived cytokines, are important and critical inflammatory mediators of a wide variety of disease states and conditions. The inhibition of these pro-inflammatory cytokines is of benefit in controlling, reducing and alleviating many of these disease states.

Accordingly, the present invention provides a method of treating a cytokine-mediated disease which comprises administering an effective cytokine-interfering amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof.

5 In particular, compounds of formula (I) or a pharmaceutically acceptable salt thereof are of use in the prophylaxis or therapy of any disease state in a human, or other mammal, which is exacerbated by or caused by excessive or unregulated IL-1, IL-8 or TNF production by such mammal's cell, such as, but not limited to, monocytes and/or macrophages.

Accordingly, in another aspect, this invention relates to a method of inhibiting the production of IL-1 in a mammal in need thereof which comprises administering to said mammal an effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof.

10 There are many disease states in which excessive or unregulated IL-1 production is implicated in exacerbating and/or causing the disease. These include rheumatoid arthritis, osteoarthritis, endotoxemia and/or toxic shock syndrome, other acute or chronic inflammatory disease states such as the inflammatory reaction induced by endotoxin or inflammatory bowel disease, 15 tuberculosis, atherosclerosis, muscle degeneration, multiple sclerosis, cachexia, bone resorption, psoriatic arthritis, Reiter's syndrome, rheumatoid arthritis, gout, traumatic arthritis, rubella arthritis and acute synovitis. Recent evidence also links IL-1 activity to diabetes, pancreatic β cells and Alzheimer's disease.

20 In a further aspect, this invention relates to a method of inhibiting the production of TNF in a mammal in need thereof which comprises administering to said mammal an effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof

25 Excessive or unregulated TNF production has been implicated in mediating or exacerbating a number of diseases including rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions, sepsis, septic shock, endotoxic shock, gram negative sepsis, toxic shock syndrome, adult respiratory distress syndrome, cerebral malaria, chronic pulmonary inflammatory disease, silicosis, pulmonary 30 sarcoisosis, bone resorption diseases, such as osteoporosis, reperfusion injury, graft vs. host reaction, allograft rejections, fever and myalgias due to infection, such as influenza, cachexia secondary to infection or malignancy, cachexia secondary to acquired immune deficiency syndrome (AIDS), AIDS, ARC (AIDS related complex), keloid formation, scar tissue formation, 35 Crohn's disease, ulcerative colitis and pyresis.

Compounds of formula (I) are also useful in the treatment of viral infections, where such viruses are sensitive to upregulation by TNF or will elicit TNF production *in vivo*. The viruses contemplated for treatment herein are those that produce TNF as a result of infection, or those which

are sensitive to inhibition, such as by decreased replication, directly or indirectly, by the TNF inhibiting-compounds of formula (1). Such viruses include, but are not limited to HIV-1, HIV-2 and HIV-3, Cytomegalovirus (CMV), Influenza, adenovirus and the Herpes group of viruses, such as but 5 not limited to, Herpes Zoster and Herpes Simplex. Accordingly, in a further aspect, this invention relates to a method of treating a mammal afflicted with a human immunodeficiency virus (HIV) which comprises administering to such mammal an effective TNF inhibiting amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof.

10 Compounds of formula (I) may also be used in association with the veterinary treatment of mammals, other than in humans, in need of inhibition of TNF production. TNF mediated diseases for treatment, therapeutically or prophylactically, in animals include disease states such as those noted above, but in particular viral infections. Examples of such 15 viruses include, but are not limited to, the lentivirus infections such as equine infectious anaemia virus, caprine arthritis virus, visna virus, or the maedi virus, or the retroviruses, such as feline immunodeficiency virus (FIV), bovine immunodeficiency virus, or canine immunodeficiency virus.

20 The compounds of formula (I) may also be used topically in the treatment or prophylaxis of topical disease states mediated by or exacerbated by excessive cytokine production, such as by IL-1 or TNF respectively, such as inflamed joints, eczema, psoriasis and other inflammatory skin conditions such as sunburn; inflammatory eye conditions including conjunctivitis; pyresis, pain and other conditions associated with 25 inflammation.

Compounds of formula (I) have also been shown to inhibit the production of IL-8 (Interleukin-8, NAP). Accordingly, in a further aspect, this invention relates to a method of inhibiting the production of IL-8 in a mammal in need thereof which comprises administering to said mammal an 30 effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof.

There are many disease states in which excessive or unregulated IL-8 production is implicated in exacerbating and/or causing the disease. These diseases are characterized by massive neutrophil infiltration such as, 35 psoriasis, inflammatory bowel disease, asthma, cardiac and renal reperfusion injury, adult respiratory distress syndrome, thrombosis and glomerulonephritis. All of these diseases are associated with increased IL-8 production which is responsible for the chemotaxis of neutrophils into the inflammatory site. In contrast to other inflammatory cytokines (IL-1, TNF,

and IL-6), IL-8 has the unique property of promoting neutrophil chemotaxis and activation. Therefore, the inhibition of IL-8 production would lead to a direct reduction in the neutrophil infiltration.

The compounds of formula (I) are administered in an amount sufficient to inhibit cytokine, in particular IL-1, IL-8 or TNF, production such that it is regulated down to normal levels, or in some case to subnormal levels, so as to ameliorate or prevent the disease state. Abnormal levels of IL-1, IL-8 or TNF, for instance in the context of the present invention, constitute: (i) levels of free (not cell bound) IL-1, IL-8 or TNF greater than or equal to 1 picogram per ml; (ii) any cell associated IL-1, IL-8 or TNF; or (iii) the presence of IL-1, IL-8 or TNF mRNA above basal levels in cells or tissues in which IL-1, IL-8 or TNF, respectively, is produced.

The discovery that the compounds of formula (I) are inhibitors of cytokines, specifically IL-1, IL-8 and TNF is based upon the effects of the compounds of formulas (I) on the production of the IL-1, IL-8 and TNF in *in vitro* assays which are described herein.

As used herein, the term "inhibiting the production of IL-1 (IL-8 or TNF)" refers to:

- 20 a) a decrease of excessive *in vivo* levels of the cytokine (IL-1, IL-8 or TNF) in a human to normal or sub-normal levels by inhibition of the *in vivo* release of the cytokine by all cells, including but not limited to monocytes or macrophages;
- 25 b) a down regulation, at the genomic level, of excessive *in vivo* levels of the cytokine (IL-1, IL-8 or TNF) in a human to normal or sub-normal levels;
- c) a down regulation, by inhibition of the direct synthesis of the cytokine (IL-1, IL-8 or TNF) as a posttranslational event; or
- 30 d) a down regulation, at the translational level, of excessive *in vivo* levels of the cytokine (IL-1, IL-8 or TNF) in a human to normal or sub-normal levels.

As used herein, the term "TNF mediated disease or disease state" refers to any and all disease states in which TNF plays a role, either by production of TNF itself, or by TNF causing another monokine to be released, such as but not limited to IL-1, IL-6 or IL-8. A disease state in which, for instance, IL-1 is a major component, and whose production or action, is exacerbated or secreted in response to TNF, would therefore be considered a disease state mediated by TNF.

As used herein, the term "cytokine" refers to any secreted polypeptide that affects the functions of cells and is a molecule which modulates interactions between cells in the immune, inflammatory or hematopoietic response. A cytokine includes, but is not limited to, monokines and lymphokines, regardless of which cells produce them. For instance, a monokine is generally referred to as being produced and secreted by a mononuclear cell, such as a macrophage and/or monocyte. Many other cells however also produce monokines, such as natural killer cells, fibroblasts, basophils, neutrophils, endothelial cells, brain astrocytes, bone marrow stromal cells, epidermal keratinocytes and B-lymphocytes. Lymphokines are generally referred to as being produced by lymphocyte cells. Examples of cytokines include, but are not limited to, Interleukin-1 (IL-1), Interleukin-6 (IL-6), Interleukin-8 (IL-8), Tumor Necrosis Factor-alpha (TNF- α) and Tumor Necrosis Factor beta (TNF- β).

As used herein, the term "cytokine interfering" or "cytokine suppressive amount" refers to an effective amount of a compound of formula (I) which will cause a decrease in the *in vivo* levels of the cytokine to normal or sub-normal levels, when given to a patient for the prophylaxis or treatment of a disease state which is exacerbated by, or caused by, excessive or unregulated cytokine production.

As used herein, the cytokine referred to in the phrase "inhibition of a cytokine, for use in the treatment of a HIV-infected human" is a cytokine which is implicated in (a) the initiation and/or maintenance of T cell activation and/or activated T cell-mediated HIV gene expression and/or replication and/or (b) any cytokine-mediated disease associated problem such as cachexia or muscle degeneration.

As TNF- β (also known as lymphotoxin) has close structural homology with TNF- α (also known as cachectin) and since each induces similar biologic responses and binds to the same cellular receptor, both TNF- α and TNF- β are inhibited by the compounds of the present invention and thus are herein referred to collectively as "TNF" unless specifically delineated otherwise.

In order to use a compound of formula (I) or a pharmaceutically acceptable salt thereof in therapy, it will normally be formulated into a pharmaceutical composition in accordance with standard pharmaceutical practice. This invention, therefore, also relates to a pharmaceutical composition comprising an effective, non-toxic amount of a compound of formula (I) and a pharmaceutically acceptable carrier or diluent.

Compounds of formula (I), pharmaceutically acceptable salts thereof and pharmaceutical compositions incorporating such may conveniently be administered by any of the routes conventionally used for drug administration, for instance, orally, topically, parenterally or by inhalation.

5 The compounds of formula (I) may be administered in conventional dosage forms prepared by combining a compound of formula (I) with standard pharmaceutical carriers according to conventional procedures. The compounds of formula (I) may also be administered in conventional dosages in combination with a known, second therapeutically active compound.

10 These procedures may involve mixing, granulating and compressing or dissolving the ingredients as appropriate to the desired preparation. It will be appreciated that the form and character of the pharmaceutically acceptable character or diluent is dictated by the amount of active ingredient with which it is to be combined, the route of administration and

15 other well-known variables. The carrier(s) must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

The pharmaceutical carrier employed may be, for example, either a solid or liquid. Exemplary of solid carriers are lactose, terra alba, sucrose, 20 talc, gelatin, agar, pectin, acacia, magnesium stearate, stearic acid and the like. Exemplary of liquid carriers are syrup, peanut oil, olive oil, water and the like. Similarly, the carrier or diluent may include time delay material well known to the art, such as glyceryl mono-stearate or glyceryl distearate alone or with a wax.

25 A wide variety of pharmaceutical forms can be employed. Thus, if a solid carrier is used, the preparation can be tableted, placed in a hard gelatin capsule in powder or pellet form or in the form of a troche or lozenge. The amount of solid carrier will vary widely but preferably will be from about 25mg. to about 1g. When a liquid carrier is used, the preparation will 30 be in the form of a syrup, emulsion, soft gelatin capsule, sterile injectable liquid such as an ampule or nonaqueous liquid suspension.

Compounds of formula (I) may be administered topically, that is by non-systemic administration. This includes the application of a compound of formula (I) externally to the epidermis or the buccal cavity and the 35 instillation of such a compound into the ear, eye and nose, such that the compound does not significantly enter the blood stream. In contrast, systemic administration refers to oral, intravenous, intraperitoneal and intramuscular administration.

Formulations suitable for topical administration include liquid or semi-liquid preparations suitable for penetration through the skin to the site of inflammation such as liniments, lotions, creams, ointments or pastes, and drops suitable for administration to the eye, ear or nose. The active 5 ingredient may comprise, for topical administration, from 0.001% to 10% w/w, for instance from 1% to 2% by weight of the formulation. It may however comprise as much as 10% w/w but preferably will comprise less than 5% w/w, more preferably from 0.1% to 1% w/w of the formulation.

Lotions according to the present invention include those suitable for 10 application to the skin or eye. An eye lotion may comprise a sterile aqueous solution optionally containing a bactericide and may be prepared by methods similar to those for the preparation of drops. Lotions or liniments for application to the skin may also include an agent to hasten drying and to cool the skin, such as an alcohol or acetone, and/or a moisturizer such as 15 glycerol or an oil such as castor oil or arachis oil.

Creams, ointments or pastes according to the present invention are semi-solid formulations of the active ingredient for external application. They may be made by mixing the active ingredient in finely-divided or powdered form, alone or in solution or suspension in an aqueous or non- 20 aqueous fluid, with the aid of suitable machinery, with a greasy or non-greasy base. The base may comprise hydrocarbons such as hard, soft or liquid paraffin, glycerol, beeswax, a metallic soap; a mucilage; an oil of natural origin such as almond, corn, arachis, castor or olive oil; wool fat or its derivatives or a fatty acid such as steric or oleic acid together with an 25 alcohol such as propylene glycol or a macrogel. The formulation may incorporate any suitable surface active agent such as an anionic, cationic or non-ionic surfactant such as a sorbitan ester or a polyoxyethylene derivative thereof. Suspending agents such as natural gums, cellulose derivatives or inorganic materials such as siliceous silicas, and other ingredients such as 30 lanolin, may also be included.

Drops according to the present invention may comprise sterile aqueous or oily solutions or suspensions and may be prepared by dissolving the active ingredient in a suitable aqueous solution of a bactericidal and/or fungicidal agent and/or any other suitable preservative, and preferably 35 including a surface active agent. The resulting solution may then be clarified by filtration, transferred to a suitable container which is then sealed and sterilized by autoclaving or maintaining at 98-100°C. for half an hour. Alternatively, the solution may be sterilized by filtration and transferred to the container by an aseptic technique. Examples of

bactericidal and fungicidal agents suitable for inclusion in the drops are phenylmercuric nitrate or acetate (0.002%), benzalkonium chloride (0.01%) and chlorhexidine acetate (0.01%). Suitable solvents for the preparation of an oily solution include glycerol, diluted alcohol and propylene glycol.

5 Compounds of formula (I) may be administered parenterally, that is by intravenous, intramuscular, subcutaneous intranasal, intrarectal, intravaginal or intraperitoneal administration. The subcutaneous and intramuscular forms of parenteral administration are generally preferred.

10 Appropriate dosage forms for such administration may be prepared by conventional techniques. Compounds of formula (I) may also be administered by inhalation, that is by intranasal and oral inhalation administration. Appropriate dosage forms for such administration, such as an aerosol formulation or a metered dose inhaler, may be prepared by conventional techniques.

15 For all methods of use disclosed herein for the compounds of formula (I), the daily oral dosage regimen will preferably be from about 0.1 to about 80 mg/kg of total body weight, preferably from about 0.2 to 30 mg/kg, more preferably from about 0.5 mg to 15 mg. The daily parenteral dosage regimen about 0.1 to about 80 mg/kg of total body weight, preferably from about 0.2 to about 30 mg/kg, and more preferably from about 0.5 mg to 15 mg/kg. The daily topical dosage regimen will preferably be from 0.1 mg to 150 mg, administered one to four, preferably two or three times daily. The daily inhalation dosage regimen will preferably be from about 0.01 mg/kg to about 1 mg/kg per day. It will also be recognized by one of skill in the art that the optimal quantity and spacing of individual dosages of a compound of formula (I) or a pharmaceutically acceptable salt thereof will be determined by the nature and extent of the condition being treated, the form, route and site of administration, and the particular patient being treated, and that such optimums can be determined by conventional techniques. It will also be appreciated by one of skill in the art that the optimal course of treatment, i.e., the number of doses of a compound of formula (I) or a pharmaceutically acceptable salt thereof given per day for a defined number of days, can be ascertained by those skilled in the art using conventional course of treatment determination tests.

30 The invention will now be described by reference to the following examples which are merely illustrative and are not to be construed as a limitation of the scope of the present invention.

Synthetic Examples

Example 1 - 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - To a solution of 2-(4-cyanophenyl)-4-(4-fluorophenyl)-N-1-hydroxy-5-(4-pyridyl)imidazole (4.5 g, 13.2 mmol) [See Ex. 10 below] in 5 DMF (50 mL) was added triethyl phosphite (3.4 mL, 20 mmol), and the resulting mixture was heated at 100 °C for 2 h. After cooling, the mixture was poured into H₂O, and the solid which formed was collected by filtration, washed with H₂O and dried *in vacuo* to afford the title compound (4.0 g, 89%). Recrystallization from CH₂Cl₂/MeOH gave white solid with a mp of 10 268-269 °C.

Example 2 - 1-Methyl-2-(4-methoxyphenyl)-4-phenyl-5-(4-pyridyl)-imidazole - (a) N-Methyl-N-(4-picoly)amine - To 4-picoly chloride, hydrochloride (10 g, 0.06 mol) was added methylamine (50 mL of 40% aqueous solution, 0.58 mol), and the resulting purple solution was stirred at 15 rt for 30 min, then poured into H₂O. The mixture was extracted with CH₂Cl₂ (6x), and the combined organic extracts were evaporated. The residue was filtered under reduced pressure through a silica gel column, eluting with a solvent gradient of 0-10% MeOH/CHCl₃ to provide the title compound as a light yellow oil (4.8 g, 66%): ¹H NMR (CDCl₃): δ 8.50 (dd, 2H); 7.20 (dd, 2H); 3.70 (s, 2H); 2.40 (s, 3H); 1.70 (br, 1H).

(b) 4-Methoxy-N-methyl-N-(4-picoly)benzamide - To a solution of N-methyl-N-(4-picoly)amine (0.40 g, 3.3 mmol) and triethylamine (1.5 mL, 10.8 mmol) in CH₂Cl₂ (15 mL) was added 4-methoxybenzoyl chloride (1.2 g, 7.3 mmol). The resulting mixture was stirred at rt for 15 min, and then partitioned between 2.5N NaOH and Et₂O. The organic extract was washed with saturated aqueous NaCl and dried (MgSO₄). The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with a solvent gradient of 2-4% MeOH/CHCl₃. The material that was isolated was triturated with Et₂O to 25 provide the title compound as a light yellow solid (0.18 g, 21%): ¹H NMR (CDCl₃): δ 8.60 (d, 2H); 7.43 (br d, 2H); 7.20 (br s, 2H); 6.90 (br d, 2H); 4.66 (br s, 2H); 3.80 (s, 3H); 3.00 (s, 3H).

(c) 1-Methyl-2-(4-methoxyphenyl)-4-phenyl-5-(4-pyridyl)-imidazole - To a solution of diisopropylamine (0.16 mL, 1.1 mmol) in THF 35 at -78 °C was added n-butyllithium (0.38 mL of 2.5 M solution, 0.95 mmol). To the resulting mixture was added a solution of 4-methoxy-N-methyl-N-(4-picoly)benzamide (0.16 g, 0.62 mmol) in THF. The resulting dark red solution was warmed to -40 °C and stirred for 15 min, at which time benzonitrile (0.13 mL, 1.2 mmol) was added. The mixture was warmed to rt

and stirred for 10 h. Aqueous NH₄Cl (0.5 mL) was added, and the mixture was concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 2-4% MeOH/CHCl₃. The material which was isolated was triturated with Et₂O and recrystallized from EtOAc to provide the title compound as an off-white solid (35 mg, 17%): mp 193-194 °C.

Example 3 - 2-(4-Cyanophenyl)-1-methyl-4-phenyl-5-(4-pyridyl)imidazole - (a) 4-Cyano-N-methyl-N-(4-picolyl)benzamide - The title compound was prepared using the same procedure as described in Example 2, step (b) except using 4-cyanobenzoyl chloride: ¹H NMR (CDCl₃): δ 8.49 (dd, 2H); 7.86-7.04 (m, 6H); 4.70 and 4.43 (two br s, 2H); 3.08 and 2.89 (two br s, 3H).

(b) 4-Cyano-N-[N"-α-dibenzoyl-1,4-dihydropyridyl-methylenyl]-N-methylbenzamide - To a solution of diisopropylamine (2.8 mL, 20 mmol) in THF at -78 °C was added n-butyllithium (6.7 mL of 2.5 M solution, 17 mmol). To the resulting mixture was added a solution of 4-cyano-N-methyl-N-(4-picolyl)benzamide (3.5 g, 14 mmol) in THF. The resulting dark purple solution was stirred at -78 °C for 10 min, at which time benzoyl chloride (4.1 mL, 35 mmol) was added. The mixture was warmed to room temperature over 30 min, then poured into aqueous NH₄Cl. The mixture was extracted with Et₂O, and the organic extract was evaporated under reduced pressure. The residue was triturated with Et₂O to provide an orange solid which was washed sparingly with acetone and copiously with Et₂O. The title compound was obtained as a bright yellow solid (1.6 g, 25%): ¹H NMR (CDCl₃): δ 7.81-7.09 (m, 16H); 6.49 (m, 2H); 3.32 (s, 3H).

(c) 2-(4-Cyanophenyl)-1-methyl-4-phenyl-5-(4-pyridyl)imidazole - To a solution of 4-cyano-N-[N"-α-dibenzoyl-1,4-dihydropyridylmethylene]-N-methylbenzamide (1.5 g, 3.3 mmol) in acetic acid (50 mL) was added ammonium acetate (1.5 g, 19.5 mmol). The resulting mixture was heated at reflux for 18 h, then allowed to cool and was concentrated. The residue was suspended in CH₂Cl₂ and filtered. The filtrate was evaporated and the residue was triturated with MeOH to afford the title compound as a white crystalline solid (0.72 g, 64%): mp 176-177 °C.

Example 4 - 2-(4-Aminomethylphenyl)-1-methyl-4-phenyl-5-(4-pyridyl)-imidazole - To a solution of 2-(4-cyanophenyl)-1-methyl-4-phenyl-5-(4-pyridyl)imidazole (0.20 g, 0.6 mmol) [See Ex. 3 above] in THF (10 mL) was added LiAlH₄ (0.60 mL of 1.0 M solution in THF, 0.6 mmol), and the

resulting mixture was stirred at rt for 1 h. The mixture was then poured into 2.5 N NaOH and extracted with Et₂O. The organic extract was evaporated, and the residue was purified by flash chromatography, eluting first with a solvent gradient of 0-10% MeOH/CHCl₃, followed by 1:10:90

5 NH₄OH/MeOH/CHCl₃. Trituration with ether afforded the title compound as a white solid (66 mg, 32%): CIMS (NH₃, *m/z*): 341 (M⁺⁺H).

Example 5 - 4-[1-Methyl-4-phenyl-5(4-pyridyl)-imidazol-2-yl] benzoic acid, sodium salt - A mixture of 2-(4-cyanophenyl)-1-methyl-4-phenyl-5(4-pyridyl)imidazole (0.10 g, 0.3 mmol) [See Ex. 3 above] in 6 N HCl (3 mL)

10 was heated at reflux for 24 h, then allowed to cool. The pH was adjusted to 7, and the solid which formed was collected by filtration and washed successively with H₂O, acetone and Et₂O to provide the title compound as a white solid (25 mg, 23%): CIMS (NH₃, *m/z*): 356 (M⁺⁺H).

Example 6 - 2-(4-Acetamidomethyphenyl)-1-methyl-4-phenyl-5-(4-pyridyl)imidazole - To a solution of 2-(4-aminomethylphenyl)-1-methyl-4-phenyl-5-(4-pyridyl)imidazole (30 mg, 0.09 mmol) [See Ex. 4 above] in pyridine (3 mL) was added acetic anhydride (0.30 mL, 3.18 mmol). The resulting solution was stirred at rt for 30 min, then concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 0-2% MeOH/CHCl₃. The isolated material was triturated with Et₂O to provide an off-white solid (10 mg, 28%) which was recrystallized from EtOAc to provide the title compound: mp 210-211 °C.

Example 7 - Methyl-4-[1-methyl-4-phenyl-5-(4-pyridyl)-imidazol-2-yl] benzoate - To a suspension of 4-[1-methyl-4-phenyl-5(4-pyridyl)-imidazol-2-yl] benzoic acid, sodium salt (20 mg, 0.06 mmol) [See Ex. 5 above] in CH₂Cl₂ (2 mL) was added triethylamine (24 mL, 0.17 mmol), followed by thionyl chloride (10 mL, 0.14 mmol). The reaction mixture was stirred at rt for 30 min, at which time MeOH (0.5 mL) was added. The mixture was stirred at rt for an additional 2 h and concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 0-1% MeOH/CHCl₃ and recrystallized from EtOAc to afford the title compound as an off-white crystalline solid (1.6 mg, 8%): mp 208-209 °C.

Example 8a - 4-(4-Fluorophenyl)-N-1-hydroxy-2-(4-hydroxyphenyl)-5-(4-pyridyl)imidazole - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 4-hydroxybenzaldehyde.

Example 8b - 4-(4-Fluorophenyl)-2-(4-hydroxyphenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure

as described in Example 1, except using 4-(4-fluorophenyl)-N-1-hydroxy-2-(4-hydroxyphenyl)-5-(4-pyridyl)imidazole (see Ex.8a): mp 214-215 °C.

Example 9 - 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzoic acid - A solution containing 2-(4-cyanophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (9.6 g, 28 mmol) [See Ex. 1 above] in

concentrated HCl (100 mL) was heated at reflux for 18 h. After cooling, the pH was adjusted to neutral by the addition of 50% aqueous NaOH. The solid which formed was collected by filtration and washed successively with H₂O, acetone and Et₂O. A portion of the solid (5 g) was dissolved in MeOH and filtered under reduced pressure through a pad of silica gel, eluting with a solvent gradient of 4-10 % MeOH/CHCl₃, followed by 2:20:80 H₂O/MeOH/CHCl₃. The title compound was isolated as a yellow solid, which was recrystallized from MeOH/CH₂Cl₂ (1.2 g, 30% adjusted yield): mp 289-290 °C.

Example 10 - 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-1-N-hydroxy-5-(4-pyridyl)imidazole - (a) 4-Fluoro-N-methoxy-N-methylbenzamide

- To a mixture containing methoxymethylamine hydrochloride (44 g, 0.45 mol) and triethylamine (138 mL, 0.99 mol) in CH₂Cl₂ (500 mL) at 0 °C was added over 30 min, 4-fluorobenzoyl chloride (50 mL, 0.41 mol). The resulting mixture was allowed to warm to rt and stirring was continued for 30 min, at which time the mixture was poured into H₂O and extracted with Et₂O. The organic extract was washed with saturated aqueous NaCl and dried (MgSO₄). Removal of the solvent *in vacuo* afforded the title compound (80 g, 100%), which was used without further purification: ¹H NMR (CDCl₃): δ 7.72 (dd, 2H); 7.06 (apparent t, 2H); 3.52 (s, 3H); 3.43 (s, 3H).

(b) 4-Fluoro-2-(4-pyridyl)acetophenone - A solution of lithium diisopropylamide was prepared at -78 °C in the usual manner from diisopropylamine (21 ml, 0.15 mol) and n-butyllithium (54 mL of 2.5 M solution in hexanes, 0.135 mol), and to this was added at -78 °C, 4-picoline (10 g, 0.108 mol). After stirring an additional 15 min at -78 °C., 4-fluoro-N-methoxy-N-methylbenzamide (20 g, 0.109 mol) was added, and the mixture was allowed to slowly warm to rt. The reaction mixture was poured into saturated aqueous NaCl and extracted with 4:1 THF/CH₂Cl₂, and the organic extract was dried (MgSO₄). The solvent was removed *in vacuo*, and to the oily brown residue was added Et₂O. The title compound was obtained as a brown solid (16.8 g, 72%) which was recrystallized from Et₂O/Hex: ¹H NMR (CDCl₃): δ 8.55 (d, 2H); 8.03 (dd, 2H); 7.16 (m, 4H); 4.24 (s, 2H).

(c) 4-Fluoro-2-hydroxyimino-2-(4-pyridyl)acetophenone - The title compound was prepared using the same procedure (US 3,940,486)

employed to prepare 2-hydroxyimino-2-(4-pyridyl)acetophenone, except using 4-fluoro-2-(4-pyridyl)acetophenone.

(d) 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-N-1-hydroxy-5-(4-pyridyl)imidazole - The title compound was prepared using the same procedure (US 3,940,486) employed to prepare 2-(t-butyl)-4-(phenyl)-N-1-hydroxy-5-(4-pyridyl)imidazole, except using 4-fluoro-2-hydroxyimino-2-(4-pyridyl)acetophenone and 4-cyanobenzaldehyde: ^1H NMR (CDCl_3): δ 8.27 (d, 2H); 7.94 (d, 2H); 7.72 (d, 2H); 7.35 (d, 2H); 7.30 (dd, 2H); 6.96 (t, 2H).

Example 11 - 2-(4-Aminomethylphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - To a solution of 2-(4-cyanophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (2.5 g, 7.3 mmol) [See Ex. 1 above] in THF (50 mL) was added LiAlH_4 (7.3 mL of 1 M solution in THF, 7.3 mmol), and the resulting mixture was heated at reflux for 2 h, at which time tlc analysis indicated that the reaction was incomplete. Additional LiAlH_4 (4.0 mL, 4.0 mmol) was added and heating was continued for 30 min. The mixture was allowed to cool, then poured into 2.5 N NaOH and extracted with THF. The organic extract was washed with saturated aqueous NaCl and concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with 9:1 $\text{CHCl}_3/\text{MeOH}$, followed by 90:10:1 $\text{CHCl}_3/\text{MeOH}/\text{NH}_3$. The material that was isolated was triturated with Et_2O to afford the title compound (1.5 g, 60%): mp 214-215 °C.

Example 12a - 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-N-1-hydroxy-5-(4-quinolyl)imidazole - **(a) 4-Fluoro-2-(4-quinolyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 4-methylquinoline: ^1H NMR (CDCl_3): δ 8.87 (d, 1H); 8.13 (m, 3H); 7.86 (d, 1H); 7.73 (apparent br t, 1H); 7.56 (apparent br t, 1H); 7.28 (d, 1H); 7.20 (t, 2H); 4.71 (s, 2H).

(b) 4-Fluoro-2-hydroxyimino-2-(4-quinolyl)acetophenone - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 4-fluoro-2-(4-quinolyl)acetophenone: ^1H NMR (DMSO-d_6): δ 9.00 (d, 1H); 8.15 (m, 3H); 7.78 (m, 1H); 7.61 (m, 2H); 7.50 (d, 1H); 7.42 (t, 2H).

(c) 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-N-1-hydroxy-5-(4-quinolyl)imidazole - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 4-fluoro-2-hydroxyimino-2-(4-quinolyl)acetophenone and 4-cyanobenzaldehyde: ^1H NMR (CDCl_3): δ 8.30 (d, 2H); 7.80 (d, 1H); 7.70 (two overlapping d, 3H); 7.46 (m, 2H); 7.36 (m, 1H); 7.11 (m, 2H); 7.01 (m, 1H); 6.75 (t, 2H).

Example 12b - 2-(4-Cyanophenyl)-4-(4-fluorophenyl)-5-(4-quinolyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 1, except using 2-(4-cyanophenyl)-4-(4-fluorophenyl)-N-1-hydroxy-5-(4-quinolyl)imidazole [see Ex. 12a]: mp 294-295 °C.

5 **Example 13 - 2-(3,5-Dibromo-4-hydroxyphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** - (a) **1-(4-fluorophenyl)-2-(4-pyridyl)-ethanediol** To a stirring solution of 2.0 g (11.2 mmol) 4-(t-butyldimethylsilyloxy)methyl pyridine in 8 ml of THF at -20° C was added 14.7 mmol of lithium di-*iso*-propyl amide in THF. Thirty minutes later 4-fluorobenzaldehyde (1.66 g, 13.4 mmol) was added at which point the solution was allowed to warm slowly to rt. The reaction was quenched with NH₄Cl and extracted with ether to afford the crude protected diol which following concentration was dissolved in THF and treated with 17 ml of a 1 molar solution of tetrabutylammonium fluoride in THF overnight. Standard aqueous workup afforded the crude diol which was further purified by column chromatography (hex/EtOAc) to yield 1.6 g (62%) of the titled material.

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(b) **1-(4-fluorophenyl)-2-(4-pyridyl)ethanedione** Oxidation of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanediol according to the oxalyl chloride method of Swern [J. Org. Chem., 44, p 4148, 1979)] gave the titled dione following extractive workup and recrystallization from hexanes m.p. 85-86.5°C.

(c) **2-(3,5-Dibromo-4-hydroxyphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** To a solution of 1-(4-fluorophenyl)-2-(4-pyridyl)ethanedione (0.25 g, 1.1 mmol) and 3,5-dibromo-4-hydroxybenzaldehyde (0.37 g, 1.3 mmol) in glacial acetic acid (5 mL) was added ammonium acetate (0.50 g, 6.5 mmol), and the resulting mixture was heated at reflux for 18 h. After cooling, the mixture was poured into H₂O, and the pH was adjusted to neutral by the addition of 2.5 N NaOH. The solid which formed was collected by filtration, washed with H₂O, dried *in vacuo* and purified by flash chromatography, eluting with a solvent gradient of 2-4% MeOH/CHCl₃. The title compound was obtained as a tan solid (15 mg, 3%): ESMS (*m/z*): 488 (M⁺+H).

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Example 14 - Ethyl 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)]-1H-imidazol-2-yl]-benzoate - A solution of 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzoic acid (30 mg, 0.08 mmol) [See Ex. 9 above] in 20% ethanolic HCl (5 mL) was heated at reflux for 24 h, cooled to rt and neutralized with 50 % NaOH. The residue was collected and purified by flash chromatography eluting with a solvent gradient of 4-10% MeOH/CHCl₃. Trituration with Et₂O afforded the title compound as a

white solid (3.2 g 66%). ^1H NMR (CDCl₃/MeOH-d₄): δ 8.45 (d, 2H); 8.12 (m, 4H); 7.52 (m, 4H); 7.15 (t, 2H); 4.42 (q, 2H); 1.43 (t, 3H).

Example 15 - 2-[3,5-Dimethyl-4-hydroxy(phenyl)]-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using 3,5-dimethyl-4-hydroxybenzaldehyde: ESMS (m/z): 360 (M⁺+H).

Example 16 - 4-(4-Fluorophenyl)-2-(2-hydroxyphenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using salicylaldehyde: ESMS (m/z): 332 (M⁺⁺H).

Example 17 - 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using 4-(methylthio)-benzaldehyde: ESMS (m/z): 362 (M⁺⁺H).

Example 18 - Methyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzoate - A mixture containing 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzoic acid, sodium salt (0.20 g, 0.5 mmol) [See Ex. 9 above] and concentrated HCl (10 drops) in MeOH (5 mL) was heated at reflux for 8 h. After cooling, the pH was adjusted to neutral by the addition of 2.5 N NaOH, and the solid which formed was collected by filtration, washed with H₂O and dried *in vacuo*. The title compound was obtained as a yellow solid (0.14 g, 76%) and was recrystallized from EtOAc/CH₂Cl₂: ^1H NMR (CDCl₃/MeOH-d₄): δ 8.36 (d, 2H); 8.03 (m, 4H); 7.60-7.30 (m, 4H); 7.07 (t, 2H); 3.84 (s, 3H).

Example 19 - 4-(4-Fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)-1H-imidazole - To a solution of 4-(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole (3.7 g, 9.8 mmol) [See Ex. 20 below] in 1:10 3 N HCl/H₂O (88 mL) was added a solution of KMnO₄ (1.5 g, 9.8 mmol) in H₂O (15 mL). After stirring at rt for 1 h, additional KMnO₄ (0.15 g, 0.9 mmol) was added, and stirring was continued for 15 min. The mixture was then poured into saturated aqueous Na₂SO₃ (200 mL), and the pH was adjusted to slightly acidic by the addition of 3 N HCl, then to neutral by the addition of 2.5 N NaOH. The solid which formed was collected by filtration, washed successively with H₂O and MeOH and recrystallized three times from MeOH to afford the title compound (0.63 g, 16%): mp 148-149 °C.

Example 20 - 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole - To a solution of 4-(4-fluorophenyl)-2-(4-

methylthiophenyl)-5-(4-pyridyl)-1H-imidazole (0.80 g, 2.2 mmol) [See Ex. 17 above] in glacial acetic acid (15 mL) was added a solution of $K_2S_2O_8$ (0.72 g, 2.6 mmol) in H_2O (20 mL). Additional glacial acetic acid (15 mL) was added to ensure homogeneity, and the resulting solution was stirred at rt for 18 h.

5 The mixture was then poured into H_2O , and the pH was adjusted to neutral by the addition of conc. NH_4OH . The solid which formed was collected by filtration to afford the title compound (0.65 g, 78%) as a tan solid, which was recrystallized from $MeOH$: mp 250-252 °C.

Example 21 - N,N-Dimethyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-

10 **imidazol-2-yl]benzamide** - To dimethylamino(methyl)aluminum chloride (0.60 mL of 0.67 M solution in toluene, 0.40 mmol) was added a solution of methyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzoate (50 mg, 0.13 mmol) [See Ex. 18 above] in 1,2-dichloroethane (5 mL). The resulting mixture was heated at reflux for 4 h, then allowed to cool and was 15 poured into 2.5 N $NaOH$. The mixture was extracted with $EtOAc$, and the organic extract was washed with saturated aqueous $NaCl$ and dried ($MgSO_4$). The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with a solvent gradient of 2-4% $MeOH/CHCl_3$ to afford the title compound (25 mg, 50%) as a white solid: 20 CIMS (NH_3 , *m/z*): 387 ($M^{++}H$).

Example 22 - 2-[(4-N,N-Dimethyl)aminomethylphenyl]-4-(4-

fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 11, except using N,N-dimethyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-25 yl]benzamide: CIMS ($NH_3, m/z$): 373 ($M^{++}H$).

Example 23 - 2-[4-(Dimethylamino)phenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using 4-(N,N-dimethylamino)benzaldehyde: ESMS (*m/z*): 359 ($M^{++}H$).

30 **Example 24 - 4-(4-Fluorophenyl)-2-phenyl-5-(4-pyridyl)-1H-imidazole** The title compound was prepared using the same procedure as described in Example 13, except using benzaldehyde: ESMS (*m/z*): 316 ($M^{++}H$).

Example 25 - 2-[4-(3-Dimethylaminopropoxy)phenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using 4-(3-dimethylamino-propoxy)benzaldehyde: ESMS (*m/z*): 417 ($M^{++}H$).

35 **Example 26 - 4-(4-Fluorophenyl)-2-(4-nitrophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as

described in Example 13, except using 4-nitrobenzaldehyde: ESMS (*m/z*): 359 (M⁺⁺H).

Example 27 - N,N-Dimethyl-4-[2-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzoyl-oxyacetamide - (a) Methyl benzylglycolate -

5 To a solution containing methyl glycolate (2.5 mL, 32 mmol) and trifluoromethyl-sulfonic acid (150 mL) in CH₂Cl₂ (10 mL) was added benzyl 2,2,2-trichloro-acetimidate (7.0 mL, 37 mmol). After stirring for several min, the mixture was poured into aqueous NaHCO₃ and extracted with Et₂O. The organic extract was washed with saturated aqueous NaCl, dried 10 (MgSO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 9-17% EtOAC/Hex to afford the title compound: ¹H NMR (CDCl₃): δ 7.34 (m, 5H); 4.62 (s, 2H); 4.11 (s, 2H); 3.78 (s, 3H).

(b) Benzyl-N,N-dimethylglycolamide - To dimethyl-

15 amino(methyl)aluminum chloride [prepared from dimethylamine hydrochloride (3.4 g, 42 mmol) and trimethyl aluminum (21 mL of 2 M solution, 42 mmol)] in toluene (40 mL) was added methyl benzylglycolate (3.0 g, 17 mmol). After stirring at rt for 1.5 h, the mixture was poured into 3 N HCl and extracted with Et₂O. The organic extract was washed with 20 saturated aqueous NaCl, dried (MgSO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 9-50% EtOAc/Hex. The title compound was obtained as a colorless oil (1.2 g, 37%): ¹H NMR (CDCl₃): δ 7.4-7.1 (m, 5H); 4.61 (s, 2H); 4.18 (s, 2H); 2.98 (s, 3H); 2.95 (s, 3H).

25 **(c) N,N-Dimethylglycolamide -** To a solution of benzyl-N,N-dimethylglycolamide (0.28 g, 1.5 mmol) in MeOH (5 mL) was added 10% palladium on activated carbon (0.15 g), and the resulting mixture was stirred under an atmosphere of H₂. After 1 h, the mixture was filtered through a pad of Celite, and the filtrate was concentrated under reduced 30 pressure to afford the title compound which was used without further purification: ¹H NMR (CDCl₃): δ 4.13 (s, 2H); 3.01 (s, 3H); 2.89 (s, 3H).

(d) N,N-Dimethyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzoyl-oxyacetamide - To a solution of 4-[4-(4-

fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzoic acid (0.15 g, 0.42 mmol)

35 [See Ex. 9 above] in DMF(10 ml) was added carbonyldiimidazole (0.34 g, 2.1 mmol). After stirring for 18 h at rt, N,N-dimethylglycolamide (0.43 g, 4.2 mmol) was added and stirring was continued for an additional 3h at rt. The reaction mixture was poured into H₂O, extracted with EtOAc and evaporated. The residue was purified by flash chromatography eluting with

a solvent gradient of 2% MeOH/CHCl₃ to afford the title compound: CIMS (NH₃,*m/z*): 445 (M⁺⁺H).

Example 28 - 2-(4-Aminophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - A mixture containing 4-(4-fluorophenyl)-2-(4-nitrophenyl)-5-(4-pyridyl)-1H-imidazole (2.0 g, 5.6 mmol) [See Ex. 26 above] and 10% palladium on activated carbon (0.4 g) was stirred under an atmosphere of H₂ for 4 h, then was filtered through a pad of Celite. The filtrate was concentrated under reduced pressure, and the residue was purified by flash chromatography, eluting with a solvent gradient of 1-10% MeOH/CHCl₃.

10 The title compound was obtained as a light orange solid (0.50 g, 27%): ¹H NMR (DMSO-d₆): δ 8.40 (d, 2H); 7.73 (d, 2H); 7.57 (m, 2H); 7.35 (m, 4H); 6.62 (t, 2H).

Example 29 - 4-(4-Fluorophenyl)-2-(4-methanesulfonamidophenyl)-5-(4-pyridyl)-1H-imidazole - To a suspension of 2-(4-aminophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (80 mg, 0.24 mmol) [See Ex. 28 above] and triethylamine (0.12 mL, 0.86 mmol) in CH₂Cl₂ (5 mL) was added methanesulfonyl chloride (55 mL, 0.72 mmol). After stirring at rt for 1 h, the mixture was poured into aqueous NaHCO₃ and extracted with EtOAc. The organic extract was washed with saturated aqueous NaCl, dried (MgSO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 2-10% MeOH/CHCl₃ to afford the title compound as a tan solid (35 mg, 36%): ESMS (*m/z*): 409 (M⁺⁺H).

Example 30 - 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl-sulfonamide - (a) **Ethyl 4-sulfonamidobenzoate** - A solution of 4-carboxybenzenesulfonamide (5.0 g, 0.025 mol) in 20% ethanolic HCl (20 mL) was heated at reflux for 18 h, then allowed to cool and was concentrated under reduced pressure to afford the title compound: ¹H NMR (CDCl₃): δ 8.20 (apparent d, 2H); 8.00 (apparent d, 2H); 4.88 (br s, 2H); 4.43 (q, 2H); 1.43 (t, 3H).

25 (b) **N-Methoxy-N-methyl-4-sulfonamidobenzamide** - To a solution of methoxymethylamino(methyl)aluminum chloride [prepared from methoxymethylamine hydrochloride (4.8 g, 50 mmol) and trimethyl aluminum (25 mL of 2 M solution, 50 mmol)] in toluene (50 mL) at 0 °C was added ethyl 4-sulfonamidobenzoate (3.8 g, 17 mmol). The mixture was allowed to warm to rt and stir for 3 h, then was poured into a slurry of silica gel (50 g) in CHCl₃ (200 mL). The mixture was filtered, and the filtrate was concentrated under reduced pressure. The residue was poured into H₂O,

and the solid which formed was collected by filtration, washed with H₂O and dried *in vacuo* to afford the title compound (1.7 g, 42%): ¹H NMR (CDCl₃/MeOH-d₄): δ 7.86 (d, 2H); 7.66 (d, 2H); 3.43 (s, 3H); 3.29 (s, 3H).

(c) 4-Formylbenzenesulfonamide - To a solution of N-
5 methoxy-N-methyl-4-sulfonamidobenzamide (1.0 g, 4.1 mmol) in THF (25 mL) at -78 °C was added LiAlH₄ (6.1 mL of 1 M solution in THF, 6.1 mmol). After stirring at -78 °C for 30 min, the mixture was poured into a slurry of silica gel (50 g) in CHCl₃ (200 mL). The mixture was filtered and the
10 filtrate was concentrated under reduced pressure. The residue was purified by flash chromatography, eluting with a solvent gradient of 2-10% MeOH/CHCl₃. The title compound was obtained as a white solid (0.12 g, 16%): ¹H NMR (CDCl₃/MeOH-d₄): δ 10.3 (s, 1H); 8.02 (d, 2H); 7.95 (d, 2H).

(d) 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]sulfonamide The title compound was prepared using the same
15 procedure as described in Example 13, except using 4-formylbenzenesulfonamide: ESMS (m/z): 395 (M⁺⁺H).

Example 31 - N'-Cyano-N-4-[4-(fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzylguanidine - To a suspension of 2-(4-aminomethylphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (0.10 g, 0.29 mmol) [See Ex. 11 above] in CH₃CN was added diphenyl
20 cyanocarbonimidate (83 mg, 0.35 mmol). After stirring at rt for 18 h, the solid which formed was collected by filtration and washed with CH₃CN. The solid was dissolved in MeOH saturated with NH₃ and stirred for 72 h. The mixture was concentrated under reduced pressure, and the residue was
25 purified by flash chromatography, eluting with a solvent gradient of 4-10% MeOH/CHCl₃. The title compound was isolated as a pale yellow solid (22 mg, 18%): mp 280-281 °C.

Example 32 - 2-[4-(Methanesulfonamido)methylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was
30 prepared using the same procedure as described in Example 29, except using 2-(4-aminomethylphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 11 above]: ESMS (m/z): 423 (M⁺⁺H).

Example 33 - 4-(4-Fluorophenyl)-2-(4-methoxyphenyl)-5-(4-pyridyl)-1H-imidazole - (a) **1-Cyano-1-(4-pyridyl)methyl 4-methoxybenzoate** -
35 The title compound was prepared using the same procedure Lantos, I. et al. (*J. Med. Chem.* 1984, 27, 72-75) employed to prepare 1-cyano-1(4-pyridyl)-methyl benzoate, except using 4-methoxybenzoyl chloride: ¹H NMR

(CDCl₃): δ 8.81 (d, 2H); 8.10 (d, 2H); 7.57 (d, 2H); 7.01 (d, 2H); 6.74 (s, 1H); 3.93 (s, 3H).

(b) **1-(4-Fluorophenyl)-2-(4-pyridyl)-2-oxoethyl 4-methoxybenzoate and 2-(4-fluorophenyl)-1-(4-pyridyl)-2-oxoethyl 4-methoxybenzoate**

5 **methoxybenzoate** - The title compounds were prepared using the same procedure Lantos et al. (J. Med. Chem. 1984, 27, 72-75) used to prepare 1-(4-Fluorophenyl)-2-(4-pyridyl)-2-oxoethyl benzoate and 2-(4-fluorophenyl)-1-(4-pyridyl)-2-oxoethyl benzoate, except using 1-Cyano-1-(4-pyridyl)methyl 4-methoxybenzoate: ¹H NMR (faster eluting isomer, CDCl₃): δ 8.78 (d, 2H); 10 8.03 (br d, 2H); 7.73 (d, 2H); 7.53 (dd, 2H); 7.10 (apparent t, 2H); 6.93 (overlapping s and d, 3H); 3.85 (s, 3H); ¹H NMR (slower eluting isomer, CDCl₃): δ 8.66 (d, 2H); 8.04 (m, 4H); 7.46 (d, 2H); 7.15 (apparent t, 2H); 6.95 (overlapping s and d, 3H); 3.87 (s, 3H).

(c) **4-(4-Fluorophenyl)-2-(4-methoxyphenyl)-5-(4-pyridyl)-1H-imidazole**

15 **imidazole** - To a solution containing a mixture of 1-(4-fluorophenyl)-2-(4-pyridyl)-2-oxoethyl 4-methoxybenzoate and 2-(4-fluorophenyl)-1-(4-pyridyl)-2-oxoethyl 4-methoxybenzoate (0.35 g, 0.96 mmol) in glacial acetic acid (7.5 mL) was added ammonium acetate (0.35 g, 4.5 mmol). The resulting mixture was heated at reflux for 18 h, then allowed to cool. The mixture 20 was concentrated under reduced pressure, and the residue was purified by flash chromatography, eluting with a solvent gradient of 33-60% EtOAc/Hex. The material which was isolated was recrystallized from MeOH/CH₂Cl₂ to provide the title compound (65 mg, 20%) as an off-white solid: mp 264-265 °C.

25 **Example 34 - 2-(4-Amino-3-iodophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** - To a solution of 2-(4-aminophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (50 mg, 0.15 mmol) [See Ex. 28 above] in glacial acetic acid (5 mL) was added a solution of ICl (24 mg, 0.15 mmol) in glacial acetic acid (1.5 mL). The resulting mixture was stirred at 30 rt for 1 h, then poured into saturated aqueous Na₂S₂O₅. The pH was adjusted to neutral by the addition of 2.5 N NaOH and extracted with EtOAc. The organic extract was concentrated under reduced pressure, and the residue was purified by flash chromatography, eluting with a solvent gradient of 2-10% MeOH/CHCl₃. The material that was isolated was 35 recrystallized from Et₂O/Hex to afford the title compound: ¹H NMR (CDCl₃): δ 8.42 (d, 2H); 8.18 (d, 1H); 7.68 (dd, 2H); 7.42 (m, 4H); 7.09 (t, 2H); 6.77 (d, 1H).

Example 35 - N-Benzyl-N-methyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide - The title compound was prepared using the same procedure as described in Example 21, except using benzylmethylaminodimethyl aluminium and methyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzoate [See Ex. 18 above]: mp 233-234 °C.

Example 36 - 2-[4-(N-Benzyl-N-methyl)aminomethylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 11, except using N-benzyl-N-methyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide [See Ex. 35 above]: ESMS (*m/z*): 449 (M⁺⁺H).

Example 37a - 4-(4-Fluorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-quinolyl)imidazole - (a) **4-Fluoro-2-(4-quinolyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 4-methylquinoline: ¹H NMR (CDCl₃): δ 8.87 (d, 1H); 8.10 (m, 3H); 7.88 (d, 1H); 7.74 (br t, 1H); 7.57 (br t, 1H); 7.20 (m, 3H); 4.73 (s, 2H).

(b) **4-Fluoro-2-hydroxyimino-2-(4-quinolyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 4-fluoro-2-(4-quinolyl)acetophenone.

(c) **4-(4-Fluorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-quinolyl)imidazole** - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 4-fluoro-2-hydroxyimino-2-(4-quinolyl)acetophenone and 4-(methylthio)benzaldehyde: ¹H NMR (CDCl₃): δ 8.03 (m, 1H); 7.80 (br d, 2H); 7.52 (d, 1H); 7.40-7.10 (m, 5H); 6.81 (br m, 3H); 6.61 (apparent t, 2H), 2.48 (s, 3H).

Example 37b - 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-quinolyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 1, except using 4-(4-fluorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-quinolyl)imidazole: ESMS (*m/z*): 412 (M⁺⁺H).

Example 38 - 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-quinolyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 20, except using 4-(4-fluorophenyl)-2-(4-methylthiophenyl)-5-(4-quinolyl)-1H-imidazole: ESMS (*m/z*): 428 (M⁺⁺H).

Example 39 - 4-(3-Chlorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 20, except using 4-(3-chlorophenyl)-2-(4-

methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 40 below]: ESMS (*m/z*): 394 (M⁺⁺H).

Example 40a - 4-(3-Chlorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - (a) 3-Chloro-N-methoxy-N-methylbenzamide

5 The title compound was prepared using the same procedure as described in Example 10 (a) except using 3-chlorobenzoyl chloride: ¹H NMR (CDCl₃): δ 7.69 (br s, 1H); 7.58 (br d, 1H); 7.42 (br d, 1H); 7.31 (dd, 1H); 3.55 (s, 3H); 3.34 (s, 3H).

10 (b) **3-Chloro-2-(4-pyridyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 4-picoline and 3-chloro-N-methoxy-N-methylbenzamide: ¹H NMR (CDCl₃): δ 8.60 (d, 2H); 8.00 (br s, 1H); 7.89 (br d, 1H); 7.60 (br d, 1H); 7.45 (t, 1H); 7.21 (d, 2H); 4.27 (s, 2H).

15 (c) **3-Chloro-2-hydroxyimino-2-(4-pyridyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 3-chloro-2-(4-pyridyl)acetophenone.

20 (d) **4-(3-Chlorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 3-chloro-2-hydroxyimino-2-(4-pyridyl)acetophenone and 4-methylthiobenzaldehyde: ¹H NMR (CDCl₃): δ 8.04 (d, 2H); 7.70 (d, 2H); 7.21-6.91 (m, 8H); 2.47 (s, 3H).

25 **Example 40b 4-(3-Chlorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 1, except using 4-(3-chlorophenyl)-N-1-hydroxy-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole: ESMS (*m/z*): 378 (M⁺⁺H).

30 **Example 41 - 4-(4-Fluorophenyl)-2-(4-formamidomethylphenyl)-5-(4-pyridyl)-1H-imidazole** - Formic acid (10 ml) was added to acetic anhydride (20 mL) and the mixture was heated to 50 °C for 15 min. 2-(4-Aminomethyl-phenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)imidazole (0.25 g, 0.73 mmol) [See Ex. 11] was added and the reaction mixture was heated to 50 °C for 2 h. The solvent was evaporated and the residue was purified by flash chromatography, eluting with a solvent gradient of 4-10% MeOH/CHCl₃. The title compound was isolated as a tan solid (0.15 g, 55%): ESMS (*m/z*): 373 (M⁺⁺H).

35 **Example 42 - 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzohydroxamic acid** - To a solution of O-benzyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imazol-2-yl]benzohydroxamic acid (0.10 g, 0.22 mmol) [See Ex. 43 below] in ethanol (10 mL) was added 10 % palladium on carbon. After stirring under an atmosphere of H₂ for 18 h, the reaction mixture was filtered through

celite and the solids were washed with ethanol. The combined filtrates were evaporated and the residue was recrystallized from 2-propanol to afford the title compound (0.040 g, 50%): ESMS (*m/z*): 375 (M⁺⁺H).

Example 43 - O-Benzyl-4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzohydroxamic acid - To a stirred suspension of O-benzylhydroxylamine hydrochloride (1.2 g, 7.8 mmol) in toluene (20 mL) at 0 °C was added trimethylaluminum (2.0 M in toluene, 3.9 mL, 7.8 mmol). The reaction mixture was warmed to rt and stirring was continued at this temperature for 1 h. Ethyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl] benzoate (1.0 g, 2.6 mmol) [See Ex. 14 above] was added and the reaction mixture was heated at reflux for 3 h. After cooling, the reaction was poured into 10% MeOH/CHCl₃ containing silica gel. The solids were removed by filtration and washed with 10% MeOH/CHCl₃. The combined washings were evaporated and the residue was purified by flash chromatography eluting with a solvent gradient of 1-10% MeOH/CHCl₃. Trituration with ether afforded the title compound as a white solid (0.25 g, 21%): ¹H NMR (CDCl₃/ MeOH-d₄): δ 8.16 (d, 2H); 7.77 (d, 2H); 7.53 (d, 2H); 7.23 (m, 5H); 7.10 (m, 4H); 6.88 (t, 2H).

Example 44 - 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide oxime - To a mixture of 2-(4-cyanophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (3.0 g, 8.7 mmol) [See Ex. 1 above] and K₂CO₃ (2.4 g, 17 mmol) in EtOH (120 mL) and H₂O (6 mL) was added hydroxylamine hydrochloride (1.2 g, 17 mmol). After heating at reflux for 24 h, the reaction mixture was poured into H₂O. The precipitate was collected, washed with H₂O and air-dried. The crude product was dissolved in acetone, silica gel was added and the solvent was evaporated. The impregnated silica gel was added to the top of a flash column and the column was eluted with a solvent gradient of 2-10% MeOH/CHCl₃ to afford the title compound as a white solid (3.0 g, 91%): ESMS (*m/z*): 374 (M⁺⁺H).

Example 45 - N¹-Methyl-N²-cyano-N-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzylguanidine - To a suspension of 2-(4-aminomethylphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (2.5 g, 7.3 mmol) [See Ex. 11 above] in CH₃CN (250 mL) was added diphenyl cyanocarbonimidate (8.8 g, 7.3 mmol). After stirring at rt for 18 h, the solid which formed was collected by filtration and washed with CH₃CN (2.1 g, 59%).

Without further purification, this material was added to methanol (100 mL) saturated with methylamine. The flask was stoppered and the reaction was stirred for 18 h at rt. The solvent and excess methylamine were evaporated and the residue was triturated with ether to give a brown solid which was further purified by flash chromatography eluting with a solvent gradient of 4-

10% MeOH/CHCl₃ to afford the title compound as a tan solid (0.33 g, 78%): CIMS (NH₃,*m/z*): 426 (M⁺+H).

Example 46a - N-1-Hydroxy-4-(3-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - (a) 3-Methoxy-N-

5 **methoxy-N-methylbenzamide** - The title compound was prepared using the same procedure as described in Example 10, step (a) except using m-anisoyl chloride: ¹H NMR (CDCl₃): δ 7.27 (m, 3H); 7.01 (m, 1H); 3.82 (s, 3H); 3.57 (s, 3H); 3.36 (s, 3H).

10 **(b) 3-Methoxy-2-(4-pyridyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 3-methoxy-N-methoxy-N-methylbenzamide: ESMS (*m/z*): 228.2 (M⁺+H).

15 **(c) 2-Hydroxyimino-3-methoxy-2-(4-pyridyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 3-methoxy-2-(4-pyridyl)acetophenone.

20 **(d) N-1-Hydroxy-4-(3-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 2-hydroxyimino-3-methoxy-2-(4-pyridyl)acetophenone and 4-(methylthio)-benzaldehyde: ESMS(*m/z*): 390 (M⁺+H).

25 **Example 46b - 4-(3-Methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole** - The title compound was prepared using the same procedure as described in Example 1, except using N-1-hydroxy-4-(3-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole: mp 215.0-216.0 °C.

30 **Example 47 - 4-(3-Methoxyphenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 38, except using 4-(3-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Example 46 above]: mp 167-168.5 °C.

35 **Example 48 - Morpholino-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide** - The title compound was prepared using the same procedure as described in Example 21 except using dimethylamino-(morpholino)aluminum chloride and ethyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-benzoate [See Ex. 14 above]: ESMS (*m/z*): 429 (M⁺+H).

40 **Example 49 - 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 13, except using 4-(4-fluorophenyl)-1-hydroxy-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)-imidazole [See Ex. 66 below]: ESMS (*m/z*): 376.2 (M⁺+H).

Example 50 - 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylsulfinylphenyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 20 except using 4-(4-fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)-1H-imidazole [See Ex. 49 above]: ESMS (*m/z*): 392.2 (M⁺⁺H).

Example 51a - 4-(4-Fluorophenyl)-N-1-hydroxy-5-(4-pyrimidinyl)-imidazole - (a) **4-Fluoro-2-(4-pyrimidinyl)acetophenone** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 4-methylpyrimidine.

10 (b) **4-Fluoro-2-hydroxyimino-2-(4-pyrimidinyl)acetophenone** - The title compound was prepared using the same procedure described in Example 10, step (c) except using 4-fluoro-2-(4-pyrimidinyl)acetophenone.

15 (c) **4-(4-Fluorophenyl)-N-1-hydroxy-5-(4-pyrimidinyl)imidazole** - The title compound was prepared using the same procedure described in Example 10, step (d) except using 4-fluorophenyl-2-hydroxyimino-2-(4-pyrimidinyl)acetophenone.

20 **Example 51b - 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyrimidinyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 1, except using 4-(4-fluorophenyl)-N-1-hydroxy-5-(4-pyrimidinyl)imidazole: CIMS (NH₃, *m/z*): 363 (M⁺⁺H).

25 **Example 52 - 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyrimidinyl)-1H-imidazole** - The title compound was prepared using the same procedure described in Example 20, except using 4-(4-fluorophenyl)-2-(4-methylthio)phenyl)-5-(4-pyrimidinyl)-1H-imidazole: CIMS (NH₃, *m/z*): 379 (M⁺⁺H).

30 **Example 53 - 4-(4-Fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyrimidinyl)-1H-imidazole** - To a solution of 4-(4-fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyrimidinyl)-1H-imidazole (0.10 g, 0.28 mmol) [See Ex. 51 above] was added 3-chloroperbenzoic acid (50%, 0.15 g, 0.42 mmol). After stirring at rt for 72 h, the solvent was evaporated and the residue was partitioned between EtOAc and 2.5 N NaOH. The organic phase was washed with brine, dried (MgSO₄) and evaporated. The residue was triturated with EtOAc to afford the title compound as a white solid (0.50 g, 46%). CIMS (NH₃, *m/z*): 395 (M⁺⁺H).

35 **Example 54 - 4-(4-Fluorophenyl)-2-(4-Morpholinomethylphenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 11 except using morpholino-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide: mp 242-243 °C.

Example 55 - 4-(4-Fluorophenyl)-2-(4-hydroxymethyl)-5-(4-pyridyl)-1H-imidazole - To a suspension of ethyl 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl] benzoate (1.0 g, 2.6 mmol) [See Ex. 14 above] in THF (25 mL) was added LiAlH₄ (1 M in THF, 7.8 mL, 7.8 mmol). After stirring at rt for 0.5 h, the reaction mixture was poured into 2.5 N NaOH and extracted three times with 2:1 EtOAc/CH₂Cl₂. The combined extracts were washed with brine, dried (MgSO₄) and evaporated to afford the title compound as a white solid (0.50 g, 54%).

Example 56 - 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-10-benzaldehyde - To a suspension of 4-(4-fluorophenyl)-2-(4-hydroxymethyl)-5-(4-pyridyl)-1H-imidazole (0.40 g, 1.2 mmol) [See Ex. 55 above] in CH₂Cl₂ (40 mL) was added pyridinium chlorochromate (0.30 g, 1.4 mmol) at rt. The reaction mixture was stirred at this temperature for 2 h, filtered through a pad of silica gel eluting with 2% MeOH/CHCl₃ and the filtrate evaporated. The residue was purified by flash chromatography eluting with 4% MeOH/CHCl₃ followed by recrystallization from CH₂Cl₂/MeOH to afford the title compound as a white solid (0.30 g, 7.3%).

Example 57 - 4-(2-Methoxyphenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 20 except using 4-(2-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 58 below]: CIMS (NH₃,*m/z*): 390 (M⁺⁺H).

Example 58a - N-1-Hydroxy-4-(2-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole - (a) 2-Methoxy-N-methoxy-N-methylbenzamide - The title compound was prepared using the same procedure as described in Example 10, step (a) except using o-anisoyl chloride: ¹H NMR (CDCl₃): δ 7.36 (m, 3H); 6.98 (dd, 1H); 3.84 (s, 3H); 3.56 (br s, 3H); 3.32 (br s, 3H).

(b) 2-Methoxy-2-(4-pyridyl)acetophenone - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 2-methoxy-N-methoxy-N-methylbenzamide.

(c) 2-Hydroxyimino-2-methoxy-2-(4-pyridyl)acetophenone - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 2-methoxy-2-(4-pyridyl)acetophenone.

(d) N-1-Hydroxy-4-(2-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 2-hydroxyimino-2-methoxy-2-(4-pyridyl)acetophenone and 4-(methylthio)benzaldehyde: ESMS (*m/z*): 390.0 (M⁺⁺H).

Example 58b - 4-(2-Methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 1, except using N-1-hydroxy-4-(2-methoxyphenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole: CIMS

5 (NH₃,*m/z*): 374.2 (M⁺⁺H).

Example 59 - 3-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl-5-methyl-4,5-dihydro-1,2,4-oxadiazole - A solution of 4-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide oxime (0.50 g, 1.3 mmol) [See Ex. 44 above] and acetaldehyde (25 mL) in ethanol (100 mL) and

10 H₂O (100 mL) was stirred at rt for seven days. The solvent was evaporated and the residue was purified by flash chromatography eluting with a solvent gradient of 2-4% CHCl₃/MeOH. Recrystallization from EtOAc afforded the title compound as a yellow solid (0.11 g, 21%): CIMS (NH₃, *m/z*): 400 (M⁺⁺H).

Example 60 - 3-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-

15 **yl]phenyl-5-methyl-1,2,4-oxadiazole** - To a solution of 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide oxime (0.10 g, 0.27 mmol) [See Ex. 44 above] in pyridine (10 mL) was added acetic anhydride (1.0 mL) at rt. After stirring at this temperature for 18 h, the reaction mixture was poured into H₂O, and the precipitate collected, washed with H₂O and dried *in vacuo*.

20 Without further purification, the crude o-acylamidoxime was dissolved in acetic acid (5 ml) and heated at reflux for 3 h. The solvent was evaporated, H₂O was added and the mixture was neutralized with aqueous NaHCO₃. The precipitate was collected, washed with H₂O, air-dried and purified by flash chromatography eluting with 4% MeOH/CHCl₃. Trituration with ether afforded the title compound as a white solid (0.030 g, 28%): CIMS (NH₃, *m/z*): 398 (M⁺⁺H).

Example 61 - 4-(3-Aminophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - A solution of 0.161 g (0.41 mmol) of 2-(4-methylthiophenyl)-4-(3-nitrophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 62 below] in 3.4 mL of

30 HOAc-H₂O (1:1) was treated with 1.81 mL (2.87 mmol) of 20% aqueous titanium (III) chloride in one single portion. The mixture was stirred at rt for 20 min, then made basic with 10% NaOH. The aqueous mixture was extracted with 95:5 CH₂Cl₂/MeOH. The organic extracts were washed with H₂O and saturated NaCl. Evaporation of solvent *in vacuo* afforded a yellow solid which 35 was filtered through a plug of silica gel, eluting with 90:10 CHCl₃/MeOH. The title compound was isolated as a pale yellow solid (0.129 g, 78%): CIMS (NH₃,*m/z*): 359.1 (M⁺⁺H).

Example 62a - N-1-Hydroxy-2-(4-methylthiophenyl)-4-(3-nitrophenyl)-5-(4-pyridyl)imidazole - (a) 1-(3-Nitrophenyl)-2-(4-pyridyl)ethanol - The

title compound was prepared using the same procedure as described in Example 10, step (b) except using 3-nitrobenzaldehyde: ^1H NMR (CDCl_3): δ 8.41 (d, 2H); 8.23 (s, 1H); 8.15 (d, 1H); 7.67 (d, 1H); 7.54 (t, 1H); 7.19 (d, 2H); 5.05 (t, 1H); 4.41 (s, 2H).

5 (b) **1-(3-Nitrophenyl)-2-(4-pyridyl)acetophenone** - To a solution of 1.0 mL (14.3 mmol) of DMSO in 55 mL of dry CH_2Cl_2 was added 1.82 mL (12.9 mmol) of trifluoroacetic anhydride at -78 °C. The mixture was stirred for 30 min, then a solution of 1-(3-nitrophenyl)-2-(4-pyridyl)ethanol (1.09 g, 4.46 mmol) in DMSO/ CH_2Cl_2 (3/11 mL) was added dropwise. The mixture was 10 stirred at -78 °C for 2 h, then 4.1 mL (29.4 mmol) of triethylamine was added dropwise. The ice bath was removed and the mixture was warmed to room temperature. The mixture was poured into saturated NH_4Cl and extracted with CH_2Cl_2 . The organic extracts were washed with saturated NH_4Cl and saturated NaCl , then dried over MgSO_4 . Removal of the solvent *in vacuo* 15 afforded a red oil which was purified by flash chromatography, eluting with a gradient of 0-3% $\text{MeOH}/\text{CHCl}_3$. The title compound was isolated as an orange oil (0.65 g, 60%): ^1H NMR (CDCl_3): δ 8.83 (s, 1H); 8.60 (d, 2H); 8.46 (d, 1H); 8.32 (d, 1H); 7.72 (t, 1H); 7.23 (d, 2H); 4.38 (s, 2H).

20 (c) **2-Hydroxyimino-1-(3-nitrophenyl)-2-(4-pyridyl)acetophenone**
The title compound was prepared using the same procedure as described in Example 10, step (c) except using 1-(3-nitrophenyl)-2-(4-pyridyl)acetophenone.

25 (d) **N-1-Hydroxy-2-(4-methylthiophenyl)-4-(3-nitrophenyl)-5-(4-pyridyl)imidazole** - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 2-hydroxyimino-1-(3-nitrophenyl)-2-(4-pyridyl)acetophenone and 4-(methylthio)benzaldehyde: ^1H NMR ($\text{CDCl}_3/\text{MeOH-d}_4$): δ 8.55 (d, 2H); 8.43 (m, 1H); 8.15 (dd, 1H); 8.06 (d, 2H); 7.78 (d, 1H); 7.51 (m, 1H); 7.45 (d, 2H); 7.32 (m, 2H); 2.57 (s, 3H).

30 **Example 62b - 2-(4-Methylthiophenyl)-4-(3-nitrophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 1, except using N-1-hydroxy-2-(4-methylthiophenyl)-4-(3-nitrophenyl)-5-(4-pyridyl)imidazole: CIMS (NH_3 , *m/z*): 389.1 (M^{++}H)

35 **Example 63 - 4-(3-Methanesulfonamidophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 29, except using 4-(3-aminophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 61 above]: ESMS (*m/z*): 437.0 (M^{++}H).

40 **Example 64 - 3-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl-1,2,4-oxadiazol-5(4H)-one** - To a mixture of 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide oxime (0.25 g, 0.67 mmol) [See Ex. 44

above] in CH_2Cl_2 (5.0 mL) and Et_3N (0.19 mL, 1.3 mmol) was added ethyl chloroformate (0.076 mL, 0.80 mmol) at rt. After 0.5 h at this temperature, the reaction mixture was poured into H_2O , extracted four times with CH_2Cl_2 and once with 10% $\text{MeOH}/\text{CH}_2\text{Cl}_2$. The organic extracts were combined and 5 evaporated. The residue was purified by flash chromatography eluting with a solvent gradient of 2-4% $\text{MeOH}/\text{CHCl}_3$. Trituration with ether afforded a white solid (0.22 g, 73%). A portion of this compound (0.10 g, 0.22 mmol) was dissolved in HOAc (2.5 mL) and heated to reflux for 18 h. The reaction mixture was poured into H_2O , neutralized with concentrated NH_4OH , extracted with 10 EtOAc and evaporated. The residue was triturated sparingly with cold EtOAc to afford the title compound as a yellow solid (0.020 g, 23%): CIMS (NH_3 , m/z): 400 (M^++H).

Example 65 - 4-(3-Acetamidophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same 15 procedure as described in Example 6, except using 4-(3-aminophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 61 above]: ESMS (m/z): 401 (M^++H).

Example 66 - 4-(4-Fluorophenyl)-1-N-hydroxy-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)imidazole - (a) **2-Methyl isonicotinic acid** - The 20 title compound was prepared using the same procedure as described in *Liebigs Ann. Chem.*, 1958, 613, 153: ESMS (m/z): 138.0 (M^++H).

(b) **Methyl 2-methylisonicotinate** - To an ice-cooled suspension of 1.32 g (9.62 mmol) of 2-methylisonicotinic acid in 20 mL of MeOH was added 1.47 mL (20.2 mmol) of thionyl chloride. The ice-bath was removed and the 25 reaction was stirred at rt. After 22 h, the MeOH was evaporated and the residue was taken up in H_2O . The aqueous mixture was neutralized with saturated NaHCO_3 , then extracted with Et_2O . The organic extracts were washed with saturated NaCl , dried over MgSO_4 , then filtered through a bed of celite. Evaporation of solvent *in vacuo* afforded the title compound as a yellow 30 liquid (0.89 g, 61%): ^1H NMR (CDCl_3): δ 8.66 (d, 1H); 7.72 (s, 1H); 7.64 (d, 1H); 3.98 (s, 3H); 2.64 (s, 3H).

(c) **Methyl 4-fluorophenylacetate** - The title compound was prepared using the same procedure as described in Example 66, step (b) except using 4-fluorophenylacetic acid: ^1H NMR (CDCl_3): δ 7.25 (dd, 2H); 7.02 (t, 2H); 3.71 (s, 3H); 3.61 (s, 2H).

(d) **2-(4-Fluorophenyl)-1-[2-methyl-(4-pyridyl)]ethanone** - To a freshly prepared solution of NaOMe (3.0 M in MeOH) was added a solution of methyl 2-methylisonicotinate (6.81 g, 45.1 mmol) in MeOH (10 mL). This was

followed by the dropwise addition of a solution of methyl 4-fluorophenylacetate (8.34 g, 49.6 mmol) in MeOH (10 mL). The MeOH was distilled off while heating the reaction mixture at 95 °C. After 17.5 h, the solid residue was cooled. Concentrated HCl (15 mL) was added, and the mixture was heated at reflux. After 4 h, the mixture was cooled then diluted with H₂O. The aqueous mixture was washed with Et₂O, adjusted to pH 5 with 1N NaOH, then adjusted to pH 8 with saturated NaHCO₃. The alkaline aqueous was extracted with EtOAc. The EtOAc extracts were washed with saturated NaCl, then dried over Na₂SO₄. Evaporation of solvent *in vacuo* afforded a red oil which was purified by column chromatography, eluting with a gradient of 0-3% MeOH/CHCl₃. The title compound was isolated as a red oil (1.5 g, 15%).

(e) **2-(4-Fluorophenyl)-2-hydroxyimino-1-[2-methyl-(4-pyridyl)]ethanone** - The title compound was prepared using the same procedure as described in Example 10, step (c) except using 2-(4-fluorophenyl)-1-[2-methyl-(4-pyridyl)]ethanone: ESMS (*m/z*): 259 (M⁺⁺H).

(f) **4-(4-Fluorophenyl)-1-hydroxy-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)imidazole** - The title compound was prepared using the same procedure as described in Example 10, step (d) except using 2-(4-fluorophenyl)-2-hydroxyimino-1-[2-methyl-(4-pyridyl)]ethanone and 4-(methylthio)benzaldehyde: ESMS (*m/z*): 392 (M⁺⁺H).

Example 67- 3-[4-(4-Fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]-phenyl-5,5-dimethyl-4,5-dihydro-1,2,4-oxadiazole - To a solution of 4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzamide oxime (0.25 g, 0.67 mmol) [See Ex. 44 above] in acetone (10 mL) was added pyridinium trifluoroacetate (0.39 g, 2.0 mmol). After heating at reflux for 18 h, the reaction mixture was poured into saturated aqueous NaHCO₃, extracted with EtOAc and the organic phase was evaporated. The residue was purified by flash chromatography eluting with a solvent gradient of 2-10% MeOH/CHCl₃ to afford the title compound as a white solid (0.12 g, 43 %): CIMS (NH₃, *m/z*): 414 (M⁺⁺H).

Example 68 - N-Hydroxy-N-1-[4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl]ethyl] urea

(a) **a-Methyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]benzyl alcohol** To a mixture of 2-(4-cyanophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole (1.0 g, 2.9 mmol) [See Ex. 1 above] was added MeMgBr (3 M in Et₂O, 4.0 mL, 12 mmol) at rt. The reaction mixture was heated at reflux for 1 h, poured into saturated aqueous NH₄Cl, extracted with THF and the organic phase was evaporated. The residue was dissolved in MeOH (20 mL) and NaBH₄ (1.0 g, 26 mmol) was added. After 0.5

h at rt, the solvent was evaporated and the residue was purified by flash chromatography eluting with a solvent gradient of 1-10% MeOH/CHCl₃ to afford the title compound as a white solid (0.26 g, 25%): ¹H NMR (CDCl₃/MeOH-d₄): δ 8.37 (d, 2H); 7.79 (d, 2H); 7.4-7.2 (m, 6H); 6.99 (t, 3H); 5 4.76 (q, 1H); 1.35 (d, 3H).

(b) **N-Hydroxy-N-[1-[4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl]ethyl]urea** - To a mixture of a-Methyl-4-[4-(4-fluorophenyl)-5-(4-pyridyl)imidazol-2-yl]benzyl alcohol (0.25 g, 0.70 mmol), P(Ph)₃ (0.46 g, 1.75 mmol) and N,O-bis(benzyloxycarbonyl)hydroxylamine (0.48 10 g, 1.75 mmol) in THF (15 mL) was added DEAD (0.28 mL, 1.75 mmol) at rt. The reaction mixture was stirred at this temperature for 3 h and the solvent evaporated. The residue was partially purified by flash chromatography eluting with 1% MeOH/CHCl₃. Methanol (25 mL) was added to this material and the mixture was cooled to -78 °C. Ammonia was bubbled in at this 15 temperature for 15 min. The reaction mixture was warmed slowly to rt, stoppered and stirred at rt for 2 days. The solvent was evaporated and the residue was purified by flash chromatography eluting with a solvent gradient of 1-10% MeOH/CHCl₃. The title compound was obtained as an off-white solid (0.43 g, 14%): FABMS (m/z): 418 (M⁺⁺H).

20 **Example 69 - N-Hydroxy-N-[4-[4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazol-2-yl]phenyl]methyl urea** - The title compound was obtained using the same procedure described in Example 68, except using 4-(4-fluorophenyl)-2-(4-hydroxymethyl)-5-(4-pyridyl)-1H-imidazole: FABMS (m/z): 418 (M⁺⁺H).

Example 70 - 4-(3-Methylthiophenyl)-2-(4-morpholinomethylphenyl)-5-(4-pyridyl)-1H-imidazole -

25 (a) **3-Methylthiobenzaldehyde** - The title compound was prepared using the same procedure as described by Campbell, J. R. in *J. Org. Chem.*, 1962, 27, 2207: ¹H NMR (CDCl₃): δ 9.95 (s, 1H); 7.72 (s, 1H); 7.61 (d, 1H); 7.45 (m, 2H); 2.53 (s, 3H).

30 (b) **1-(3-Methylthiophenyl)-2-(4-pyridyl)ethanol** - The title compound was prepared using the same procedure as described in Example 10, step (b) except using 3-(methylthio)benzaldehyde: ¹H NMR (CDCl₃): δ 8.33 (d, 2H); 7.0-7.5 (m, 6H), 4.87 (m, 1H); 2.96 (m, 2H); 2.45 (s, 3H).

35 (c) **1-(3-Methylthiophenyl)-2-(4-pyridyl)ethanedione** - To a solution of 1-(3-methylthiophenyl)-2-(4-pyridyl)ethanol (2.5 g, 10.2 mmol) in CH₂Cl₂ (150 mL) was added a mixture of celite (4.4 g) and pyridinium dichromate (4.4 g, 20.4 mmol). After stirring for 12 h, the mixture was filtered through celite. The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with a solvent gradient of 40-50% EtOAc/Hex to

provide the title compound (144 mg, 5.5%): ^1H NMR (CDCl_3): δ 8.88 (br d, 2H); 7.85 (s, 1H); 7.78 (d, 2H); 7.67 (d, 1H); 7.56 (d, 1H); 7.44 (t, 1H); 2.55 (s, 3H).

5 **(d) 4-Morpholinomethylbenzaldehyde diethyl acetal** - The title compound was prepared using the same procedure as described by Borch, R. F., Bernstein, M. D., and Durst, H. D. in *J. Am. Chem. Soc.*, 1971, 93, 2897 except using the diethyl acetal: ^1H NMR (CDCl_3): δ 7.41 (d, 2H); 7.32 (d, 2H,); 5.48 (s, 1H); 3.3-3.8 (m, 10H); 2.43 (br s, 4H); 1.25 (t, 6H).

10 **(e) 4-(3-Methylthiophenyl)2-(4-morpholinomethylphenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 13 except using 1-(3-methylthiophenyl)-2-(4-pyridyl)-ethanedione and 4-morpholinomethylbenzaldehyde diethyl acetal: ^1H NMR (CDCl_3): δ 8.47 (d, 2H); 8.02 (d, 2H); 7.3-7.9 (m, 8H); 3.72 (t, 4H); 3.54 (s, 2H); 2.44 (br s, 4H); 2.38 (s, 3H).

15 **Example 71 - 4-(3-Methylsulfinylphenyl)-2-(4-morpholinomethylphenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described for Example 20, except using 4-(3-methylthiophenyl)2-(4-morpholinomethylphenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 70 above]: ^1H NMR (CDCl_3): δ 8.38 (d, 2H); 7.92 (d, 2H); 7.1-7.6 (m, 8H); 3.76 (t, 4H); 3.59, (s, 2H); 2.73 (s, 3H,); 2.52 (br s, 4H).

20 **Example 72 - 4-(3-Methanesulfonamidophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 20, except 25 using 4-(3-methanesulfonamido-phenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 63 above]: CIMS ($\text{NH}_3, m/z$): 453.3 (M^++H).

25 **Example 73 - 2-(4-Ethylthiophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 13, except using 4-ethylthiobenzaldehyde: mp 203-30 205 °C.

30 **Example 74 - 2-(4-Ethylsulfinylphenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 20, except using 2-(4-ethylthiophenyl)-4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole [See Ex. 73 above]: mp 240 °C.

35 **Example 75 - 4-(4-Fluorophenyl)-2-[4-(4-methyl-1-piperazinyl)-sulfonyl-phenyl]-5-(4-pyridyl)-1H-imidazole**

(a) **Ethyl [4-(4-methyl piperazinyl) sulfonamido] benzoate** - A mixture of 4-chlorosulfonyl benzoic acid (5.0 g, 22.67 mmol), N-methyl

piperazine (25 mL) and MeOH (5 mL) was stirred for 18 h and ether (200 mL) was added to the mixture. The crystalline solid precipitate was filtered and washed with ether (200 mL). The solid was suspended in 20% ethanolic HCl and the mixture was heated at reflux until a homogeneous solution was 5 attained (about 2 h). The solution was cooled to rt, concentrated, and the residue was partitioned between sat. NaHCO₃ and EtOAc. The organic extract was dried and concentrated to yield the title compound (5.8 g 80%).

10 (b) **4-(4-Methyl piperazinyl) sulfonamido benzyl alcohol** - The title compound was prepared using the same procedure as described in Example 78, step (b) except using ethyl [4-(4-methyl piperazinyl) sulfonamido] benzoate.

15 (c) **4-(4-Methyl piperazinyl) sulfonamido benzaldehyde** - To a solution of oxalyl chloride (1.06 mL, 12.1 mmol) in CH₂Cl₂ (20 mL) was added DMSO (1.8 mL, 25.4 mmol.) at -60 °C and the mixture was stir red for 25 min. A solution of 4-(4-methyl piperazinyl) sulfonamido benzyl alcohol (3.0 g, 10.5 mmol) in CH₂Cl₂ (25 mL) and DMSO (5 mL) was added. and the mixture was stirred for 1.5 h at -60 °C. Triethylamine (7.4 mL) was added and the mixture was partitioned between brine and EtOAc. The organic extract was concentrated, then purified by flash chromatography to 20 yield the title compound (1.0 g, 33%).

25 (d) **4-(4-Fluorophenyl)-2-[4-(4-methyl piperazinyl) sulfonamido phenyl]-5-(4-pyridyl)-1H-imidazole** - The title compound was prepared using the same procedure as described in Example 13, except using 4-(4-methyl piperazinyl) sulfonamido benzaldehyde: mp 74-76 °C.

20 **Example 76 - 4-(4-Fluorophenyl)-2-[4-(N-methylmethanesulfonamido)-methylphenyl]-5-(4-pyridyl)-1H-imidazole**

30 (a) **Methyl 4-[(methanesulfonamido)methyl]benzoate** - To a suspension of 4-(aminomethyl)benzoic acid (10 g, 66 mmol) in MeOH (100 mL) at 0 °C was added SOCl₂ (5.3 mL, 73 mmol) dropwise. The ice bath was removed and the reaction stirred at rt overnight. After heating the reaction at reflux for 4 h, the solvent was evaporated. The residue was suspended in CH₂Cl₂ (100 mL) at 0 °C and triethylamine (25 mL) was added, followed by the dropwise addition of methanesulfonyl chloride (7.75 mL, 100 mmol). The reaction was stirred at rt for 1 h, poured into ice H₂O, extracted with 35 CH₂Cl₂, dried over anhydrous Na₂SO₄, filtered and evaporated. The crude product was flash chromatographed on silica gel eluting with 1% MeOH/CHCl₃. The title compound was isolated as a white solid (11.8 g, 74 %): ¹H NMR (CDCl₃): δ 8.03 (d, 2H); 7.42 (d, 2H); 4.9 (br t, 1H); 4.38 (d, 2H); 3.92 (s, 3H); 2.89 (s, 3H).

(b) Methyl 4-[(N-Methylmethanesulfonamido)methyl]benzoate

- To a mixture of methyl 4-[(methanesulfonamido)methyl]benzoate (5 g, 20.6 mmol) in MeOH (100 mL) at rt was added K₂CO₃ (2.9 g, 21 mmol). Methyl iodide (7 ml, 16 g, 112 mmol) was added and the mixture stirred overnight.

5 The reaction was filtered and the solid washed with CHCl₃/MeOH. The combined filtrates were evaporated and the residue was purified by flash chromatography on eluting with 0-5% MeOH/ CHCl₃. The title compound was isolated as a white solid (4.9 g, 94%): ¹H (CDCl₃): δ 8.08 (d, 2H); 7.48 (d, 2H); 4.41 (s, 1H); 3.97 (s, 3H); 3.91 (s, 3H); 2.83 (s, 3H).

10 (c) 4-[(N-Methylmethanesulfonamido)methyl]benzyl alcohol

The title compound was prepared using the same procedure as described in Example 78, step (b) except using methyl 4-[(N-methylmethane-sulfonamido)methyl]benzoate: ¹H NMR (CDCl₃): δ 7.34 (m 4H); 4.68 (s, 2H); 4.29 (s, 2H); 2.83 (s, 3H); 2.74 (s, 3H).

15 (d) 4-[(N-Methylmethanesulfonamido)methyl]benzaldehyde

The title compound was prepared using the same procedure as described in Example 78, step (c) except using 4-[(N-methylmethanesulfonamido)methyl]benzyl alcohol: ¹H NMR (CDCl₃): δ 10.02 (s, 1H); 7.9 (d, 2H); 7.54 (d, 2H); 4.4 (s, 2H); 2.9 (s, 3H); 2.81 (s, 3H).

20 (e) 4-(4-Fluorophenyl)-2-[4-(N-methylmethanesulfonamido)methylphenyl]-5-(4-pyridyl)-1H-imidazole - The title compound was prepared using the same procedure as described in Example 13, except using 4-[(N-methylmethane-sulfonamido)methyl]benzaldehyde: mp 222-224 °C.

Example 77 - Diethyl [1-methyl-4-phenyl-5-(4-pyridyl)-imidazol-2-

25 yl]methoxy]methylphosphonate

(a) **N-Methyl-N-[4-picollyl]formamide** - To a solution of 4-picollyl chloride-HCl (15 g, 91.4 mmol) and N-methylformamide (53.4 mL, 914 mmol) in 300 mL of THF was added 80% NaH in mineral oil (5.48 g, 183 mmol). After stirring at rt for 18 h the mixture was quenched with ice water and partitioned between CH₂Cl₂ and H₂O. The organic extract was washed with aqueous NaCl and dried over MgSO₄. The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with 50:1 CH₂Cl₂/MeOH. The title compound was obtained as a pale yellow oil (10.5 g, 76%): ESMS (m/z): 151 (M⁺⁺H).

30 (b) **1-Methyl-4-phenyl-5-[4-pyridyl]imidazole** - To a solution of di-*iso*-propyl-amine (11.2 mL, 79.9 mmol) in 150 mL of THF at -78 °C was added n-butyllithium (31.9 mL of 2.5 M solution, 79.9 mmol). To the resulting mixture was added a solution of N-methyl-N-[4-picollyl]formamide (10 g, 66.5 mmol) in THF. The resulting orange-brown solution was stirred

at -78 °C for 20 min , at which time benzonitrile (13.6 mL, 133 mmol) was added. The resulting dark brown mixture was allowed to warm to rt and stirred for 1 h, heated to reflux for 4 h, and then cooled to rt and partitioned between CH₂Cl₂ and H₂O. The organic extract was washed with aqueous 5 NaCl and dried (MgSO₄). The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with 50:1 CH₂Cl₂/MeOH. The title compound was obtained as a light tan solid (5.83 g, 37%): mp 158-159 °C.

(c) **2-Formyl-1-methyl-4-phenyl-5-[4-pyridyl]imidazole** - To a 10 solution of 1-methyl-4-phenyl-5-[4-pyridinyl]imidazole (0.275 g, 1.17 mmol) in THF at -78 °C was added n-butyllithium (0.56 mL of 2.5 M solution, 1.40 mmol). The resulting red-orange solution was allowed to stir at -78 °C for 0.5 h when DMF (0.18 mL, 2.34 mmol) was added. The mixture was allowed to warm to rt and stir for 4 h, then quenched with ice water and partitioned between 15 CH₂Cl₂ and H₂O. The organic extract was washed with aqueous NaCl and dried (MgSO₄). The solvent was removed *in vacuo* and the residue was purified by flash chromatography eluting with 50:1 CH₂Cl₂/MeOH. The title compound was obtained as a white solid (0.187 g, 61%): mp 167-168 °C.

(d) **2-Hydroxymethyl-1-methyl-4-phenyl-5-(4-pyridyl)imidazole** - 20 To a solution of 2-formyl-1-methyl-4-phenyl-5-[4-pyridyl] imidazole (0.830 g, 3.15 mmol) in MeOH at 0 °C was added NaBH₄ (0.143 g, 3.78 mmol). The mixture was stirred at rt for 0.5 h when the solvent was evaporated *in vacuo* and the residue was partitioned between CH₂Cl₂ and H₂O. The organic extract was washed with aqueous NaCl and dried (MgSO₄). The solvent was removed 25 *in vacuo* and the residue was purified by flash chromatography eluting with 25:1 CH₂Cl₂/MeOH. The title compound was obtained as a white solid (0.608 g, 73%): mp 236-238 °C.

(e) **Diethyl [1-methyl-4-phenyl-5-(4-pyridyl)-imidazol-2-yl]methoxy-methyl-phosphonate** - To a suspension of 80% NaH in mineral 30 oil (0.013 g, 0.452 mmol) in DMF at 0 °C was added 1-methyl-2-hydroxymethyl-4-phenyl-5-[4-pyridinyl] imidazole (0.100 g, 0.377 mmol) in DMF. The resulting bright yellow solution was stirred at 0 °C for 0.5 h when diethyl chloromethylphosphonate (0.070 mL, 0.452 mmol) dissolved in 0.079 mL of HMPA was added. The resulting mixture was stirred at 0 °C for 15 min 35 and then warmed to rt. After 5 h, the solution was partitioned between CH₂Cl₂ and H₂O. The organic extract was washed with aqueous NaCl and dried (MgSO₄). The solvent was removed *in vacuo*, and the residue was purified by flash chromatography, eluting with 50:1 CH₂Cl₂/MeOH. The title compound was obtained as a light amber oil (0.088 g, 56%): ESMS (*m/z*): 416 (M⁺+H).

Example 78- 4-(4-Fluorophenyl)-2-[4-(methanesulfonamido)-methylphenyl]-5-(4-pyridyl)-1H-imidazole - (a) Methyl 4-[(methanesulfonamido)methyl]-benzoate - The title compound was prepared using the same procedure as described in Example 76, step (a).

5 **(b) 4-[(Methanesulfonamido)methyl]benzyl alcohol** - To a mixture of methyl 4-[(methanesulfonamido)methyl]benzoate (3.6 g, 15 mmol) in THF (150 mL) was added LiAlH₄ (1 M in THF, 30 mL, 30 mmol). The reaction mixture was stirred at rt for 1 h and poured into 10% MeOH/CHCl₃ containing silica gel. The solids were removed by filtration, washed with 10% MeOH/CHCl₃ and the combined washings were evaporated to yield the title compound as a white solid (2.6 g, 80%).

10 **(c) 4-[(Methanesulfonamido)methyl]benzaldehyde** - To a solution of 4-[(methanesulfonamido)methyl]benzyl alcohol (1.0 g, 4.6 mmol) in CH₂Cl₂ (25 mL) was added pyridinium chlorochromate (1.5 g, 7.0 mmol). The reaction 15 mixture was stirred for 1 h at rt and poured through a pad of silica gel eluting with 2% MeOH/CHCl₃. The title compound was isolated as a tan solid (1.0 g, 100 %): ¹H NMR (CDCl₃): δ 10.03 (s, 1H); 7.88 (d, 2H); 7.57 (d, 2H); 4.79 (br s, 1H); 4.43 (d, 2H); 2.93 (s, 3H).

15 **(d) 4-(4-Fluorophenyl)-2-[4-(methanesulfonamido)methylphenyl]-5-(4-pyridyl)-1H-imidazole** - The title compound [also prepared in Example 32] was prepared using the same procedure as described in Example 13, except using 4-[(methanesulfonamido)methyl]benzaldehyde.

Example 79 - 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole

20 **(a) 1-(t-Butyldimethylsilyloxy)-2-(4-fluorophenyl)-1-(4-pyridyl)ethanone** - To a -20 °C solution of diisopropylamine (64.4 mL, 0.46 mol) and THF (120 mL) was added 207.8 mL (0.52 mol, 2.5 M solution in hexanes) of n-butyllithium dropwise over 15 min. The temperature was lowered to -15 °C and the mixture was stirred for 0.5 hr. The solution was cooled to -20 °C and 98.14 g (0.44 mol) of 4-(t-butyldimethylsilyloxy)methyl pyridine was added dropwise over 20 min. After stirring at -20°C for 45 min, a solution of 4-fluoro-N-methoxy-N-methylbenzamide (84.5 g, 0.46 mol) [See Ex. 10, step (a)] in THF (90 mL) was added dropwise over 0.5 hr. Once the addition was complete, the ice bath was removed and the reaction mixture was warmed to 0 °C for 1 hr, then stirred at rt for 1.5 hr. The mixture was poured into a solution of NH₄Cl (98 g) and H₂O (500 mL), then extracted with EtOAc (3 x 250 mL). The EtOAc extracts were washed with H₂O and saturated NaCl, then dried over MgSO₄. Evaporation of the solvent *in vacuo* afforded the title compound as an amber oil (114.2 g, 75%).

(b) **4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)-1H-imidazole** - To a solution of 1-(t-butyldimethylsilyloxy)-2-(4-fluorophenyl)-1-(4-pyridyl)ethanone (6.3 g, 18.3 mmol) in glacial acetic acid (125 mL) was added anhydrous copper (II) acetate (6.6 g, 36.5 mmol), ammonium acetate (14 g, 183 mmol) and 4-(methylthio)benzaldehyde (3.5 g, 22.9 mmol) and the mixture was heated at reflux. After 1 hr, the reaction was cooled then poured into a mixture of conc. NH₄OH (175 mL), ice (100 mL) and EtOAc (100 mL). The resulting mixture was stirred for 15 min, then the layers were separated. The aqueous layer was extracted with EtOAc (2 x 50 mL). The combined EtOAc extracts were washed with saturated NaCl and dried over MgSO₄. Evaporation of solvent *in vacuo* gave an oil which was taken up in acetone. 3 N HCl was added dropwise to adjust the pH to 2-3, and the resulting solid was filtered. The title compound [also prepared in Ex. 17 as the free base] was isolated as the yellow hydrochloride salt (3.7 g, 51%).

15 **Example 80 - 2-[4-[(N-Benzyl-N-methyl) aminomethyl]phenyl]4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole**

(a) **4-[(N-benzyl-N-methyl)aminomethyl]benzaldehyde diethylacetal** - To 62.4 g (0.30 mol) of terephthalaldehyde monodiethyl acetal was added 32.1 g (0.30 mol) of benzyl amine and 500 mL toluene. The 20 resulting solution was heated at reflux using a Dean-Stark trap. After 1 hour the solution was cooled and concentrated to give a light yellow oil (89.1 g). The oil was dissolved in 900 mL of EtOAc and 2.0 g of 5% palladium on charcoal was added. The mixture was hydrogenated on a Parr hydrogenation apparatus under 37 psi hydrogen pressure. The mixture was 25 shaken for 1 hour at rt. The bottle was vented and 34.4 mL (0.42 mol) of 37.5% formaldehyde solution (aqueous) was added. The bottle was repressurized with 33 psi hydrogen and the mixture was shaken for 17 hours at rt. The bottle was vented and the reaction mixture was filtered and the filtrate concentrated to a nearly colorless oil (93.9 g). Vacuum distillation 30 gave 71.4 g (76%) of 4-(N-methyl-N-benzyl)aminomethylbenzaldehyde diethylacetal: bp (30 torr) 212-234 °C.

(b) **2-[4-[(N-Benzyl-N-methyl) aminomethyl]phenyl]4-(4-fluorophenyl)-5-(4-pyridyl)-1H-imidazole** - The title compound [also prepared in Ex. 36] was prepared as described in Example 13, except using 4-[(N-benzyl-N-methyl)aminomethyl]benzaldehyde diethylacetal.

BIOLOGICAL EXAMPLES

The cytokine-inhibiting effects of compounds of the present invention were determined by the following *in vitro* assays:

1. **IL-1** - Human peripheral blood monocytes were isolated and purified from either fresh blood preparations from volunteer donors, or from blood bank buffy coats, according to the procedure of Colotta *et al*, *J Immunol*, **132**, 936 (1984). These monocytes (1×10^6) were plated in 24-well plates at a concentration of 1-2 million/ml per well. The cells were allowed to adhere for 2 hours, after which time non-adherent cells were removed by gentle washing. Test compounds were then added to the cells for 1h before the addition of lipopolysaccharide (50 ng/ml), and the cultures were incubated at 37°C for an additional 24h. At the end of this period, culture supernatants were removed and clarified of cells and all debris. Culture supernatants were then immediately assayed for IL-1 biological activity, either by the method of Simon *et al.*, *J. Immunol. Methods*, **84**, 85, (1985) (based on ability of IL-1 to stimulate a Interleukin 2 producing cell line (EL-4) to secrete IL-2, in concert with A23187 ionophore) or the method of Lee *et al.*, *J. ImmunoTherapy*, **6** (1), 1-12 (1990) (ELISA assay). Compounds of formula (I) were shown to be inhibitors of *in vitro* IL-1 produced by human monocytes.
2. **TNF** - Human peripheral blood monocytes were isolated and purified from either blood bank buffy coats or plateletpheresis residues, according to the procedure of Colotta, R. *et al.*, *J Immunol*, **132**(2), 936 (1984). The monocytes were plated at a density of 1×10^6 cells/ml medium/well in 24-well multi-dishes. The cells were allowed to adhere for 1 hour after which time the supernatant was aspirated and fresh medium (1ml, RPMI-1640, Whitaker Biomedical Products, Whitaker, CA) containing 1% fetal calf serum plus penicillin and streptomycin (10 units/ml) added. The cells were incubated for 45 minutes in the presence or absence of a test compound at 1nM-10μM dose ranges (compounds were solubilized in dimethyl sulfoxide/ethanol, such that the final solvent concentration in the culture medium was 0.5% dimethyl sulfoxide/0.5% ethanol). Bacterial lipopolysaccharide (*E. coli* 055:B5 [LPS] from Sigma Chemicals Co.) was then added (100 ng/ml in 10 ml phosphate buffered saline) and cultures incubated for 16-18 hours at 37°C in a 5% CO₂ incubator. At the end of the incubation period, culture supernatants were removed from the cells, centrifuged at 3000 rpm to remove cell debris. The supernatant was then assayed for TNF activity using either a radio-immuno or an ELISA assay, as described in WO 92/10190 and by Becker *et al.*, *J Immunol*, **147**, 4307. Compounds of formula (I) were shown to be inhibitors of *in vitro* TNF production.

IL-1 and TNF inhibitory activity does not seem to correlate with the property of the compounds of Formula (I) in mediating arachidonic acid

metabolism inhibition, further the ability to inhibit production of prostaglandin and/or leukotriene synthesis, by nonsteroidal anti-inflammatory drugs with potent cyclooxygenase and/or lipoxygenase inhibitory activity does not mean that the compound will necessarily also

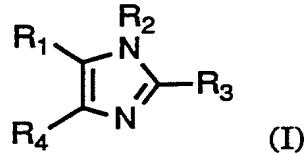
5 inhibit TNF or IL-1 production, at non-toxic doses.

3. **IL-8** - Primary human umbilical cord endothelial cells (HUVEC) (Cell Systems, Kirland, Wa) were maintained in culture medium supplemented with 15% fetal bovine serum and 1% CS-HBGF consisting of aFGF and heparin. The cells were then diluted 20-fold before being plated (250 μ l) into

10 gelating coated 96-well plates. Prior to use, culture medium was replaced with fresh medium (200 μ l). Buffer or test compound (25 μ l, at concentrations between 1 and 10 μ M) was then added to each well in quadruplicate wells and the plates incubated for 6h in a humidified incubator at 37°C in an atmosphere of 5% CO₂. At the end of the incubation period, supernatant

15 was removed and assayed for IL-8 concentration using an IL-8 ELISA kit obtained from R&D Systems (Minneapolis, MN). All data were presented as mean value (ng/ml) of multiple samples based on the standard curve. IC₅₀'s where appropriate were generated by non-linear regression analysis. The compounds of formula (I), examples 5, 8b and 9, demonstrated a dose

20 dependent reduction in the production of IL-8 (a 50-65% inhibition of IL-8).

Claims**1. A compound of formula (I):**

5 wherein

R₁ is 4-pyridyl, pyrimidinyl, quinolyl, isoquinolinyl, 1-imidazolyl or 1-benzimidazolyl which is unsubstituted or substituted with one or two substituents each of which is independently selected from C₁-4 alkyl, halo, C₁-4 alkoxy, C₁-4 alkylthio, NH₂, mono- or di-C₁-6-alkylamino or N-heterocyclyl ring which ring has from 5 to 7 members and optionally contains an additional heteroatom selected from oxygen, sulfur or NR₂₂;

10 R₂ is R₈ or -OR₁₂;

R₃ is -X_aP(Z)(X_bR₁₃)₂ or an optionally substituted aryl or heteroaryl

15 group Q;

X_a is -NR₈- , -O- , -S- or a C₁-10 alkylene chain optionally substituted by C₁-4 alkyl and optionally interrupted by -NR₈- , -O- or -S-;

X_b is -(CR₁₀R₂₀)_n , -NR₈- , -O- or -S-;

Z is oxygen or sulfur;

20 n is 0 or an integer from 1 to 10;

R₄ is phenyl, naphth-1-yl or naphth-2-yl which is optionally substituted by one or two substituents, each of which is independently selected, and which, for a 4-phenyl, 4-naphth-1-yl or 5-naphth-2-yl substituent, is halo, cyano, C(Z)NR₇R₁₇, C(Z)OR₂₃,

25 -(CR₁₀R₂₀)_mCOR₃₆ , -SR₅ , -SOR₅ , -OR₃₆ , halo-substituted-C₁-4 alkyl, C₁-4 alkyl, -ZC(Z)R₃₆ , -NR₁₀C(Z)R₂₃ , or -(CR₁₀R₂₀)_mNR₁₀R₂₀ and which, for other positions of substitution, is halo, cyano, C(Z)NR₁₆R₂₆ , C(Z)OR₈ , -(CR₁₀R₂₀)_mCOR₈ , S(O)_mR₈ , OR₈ , halo-substituted-C₁-4 alkyl, -C₁-4 alkyl, -NR₁₀C(Z)R₈ , -NR₁₀S(O)_mR₁₁ , -ZC(Z)R₈ or -(CR₁₀R₂₀)_mNR₁₆R₂₆;

30 m is 0, 1 or 2;

R₅ is hydrogen, C₁-4 alkyl, C₂-4 alkenyl, C₂-4 alkynyl or NR₇R₁₇ , excluding -SR₅ being -SNR₇R₁₇ and -SOR₅ being -SOH or ;

R₆ is C₁-4 alkyl, halo-substituted-C₁-4 alkyl, C₂-4 alkenyl, C₂-4 alkynyl or C₃-5 cycloalkyl;

R₇ and R₁₇ is each independently selected from hydrogen or C₁₋₄ alkyl or R₇ and R₁₇ together with the nitrogen to which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or

5 NR₂₂;

R₈ is hydrogen, heterocyclyl, heterocyclalkyl or R₁₁;

R₁₀ and R₂₀ is each independently selected from hydrogen or C₁₋₄ alkyl;

R₁₁ is C₁₋₁₀ alkyl, halo-substituted C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, C₃₋₇ cycloalkyl, C₅₋₇ cycloalkenyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl;

10 R₁₂ is hydrogen, -C(Z)R₁₃ or optionally substituted C₁₋₄ alkyl, aryl or aryl-C₁₋₄ alkyl;

R₁₃ is hydrogen, C₁₋₁₀ alkyl, cycloalkyl, heterocyclyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl;

15 R₁₆ and R₂₆ is each independently selected from hydrogen or optionally substituted C₁₋₄ alkyl, aryl or aryl-C₁₋₄ alkyl, or together with the nitrogen which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or NR₁₂;

20 R₂₂ is R₁₀ or C(Z)-C₁₋₄ alkyl;

R₂₃ is C₁₋₄ alkyl, halo substituted C₁₋₄ alkyl, or C₃₋₅ cycloalkyl;

R₃₆ is hydrogen or R₂₃;

25 or a pharmaceutically acceptable salt thereof;

and excluding 2-(4-chlorophenyl)-4-(4-methoxyphenyl)-5-(4-pyridyl)imidazole.

2. A compound as claimed in claim 1 wherein R₁ is optionally substituted 4-pyridyl, 4-pyrimidinyl, 4-quinolyl, 6-isoquinolinyl, 1-imidazolyl or 1-benzimidazolyl.

30

3. A compound as claimed in claim 1 or 2 wherein R₂ is hydrogen or C₁₋₁₀ alkyl.

35 4. A compound as claimed in any one of claims 1 to 3 wherein, in R₃, the group Q comprises an optionally substituted phenyl, pyrrolyl, pyridyl or pyrimidyl moiety.

5. A compound as claimed in any one of claims 1 to 4 wherein Q is substituted by up to three substituents Y₁ each of which is independently selected from C₁₋₅ alkyl, halo-substituted C₁₋₅ alkyl, halogen, -X_a-P(Z)-(X_bR₁₃)₂ or -(CR₁₀R₂₀)_nY₂ wherein Y₂ is -OR₈,

5 -NO₂, -S(O)_m'R₁₁, -SR₈, -S(O)_m'OR₈, -S(O)_mNR₈R₉, -NR₈R₉, -O(CR₁₀R₂₀)_nNR₈R₉, -C(O)R₈, -CO₂R₈, -CO₂(CR₁₀R₂₀)_nCONR₈R₉, -ZC(O)R₈, -CN, -C(Z)NR₈R₉, -NR₁₀C(Z)R₈, -C(Z)NR₈OR₉, -NR₁₀C(Z)NR₈R₉, -NR₁₀S(O)_mR₁₁, -N(OR₂₁)C(Z)NR₈R₉, -N(OR₂₁)C(Z)R₈, -C(=NOR₂₁)R₈, -NR₁₀C(=NR₁₅)SR₁₁,

10 -NR₁₀C(=NR₁₅)NR₈R₉, -NR₁₀C(=CR₁₄R₂₄)SR₁₁, -NR₁₀C(=CR₁₄R₂₄)NR₈R₉, -NR₁₀C(O)C(O)NR₈R₉, -NR₁₀C(O)C(O)OR₁₀, -C(=NR₁₃)NR₈R₉, -C(=NOR₁₃)NR₈R₉, -C(=NR₁₃)ZR₁₁, -OC(Z)NR₈R₉, -NR₁₀S(O)_mCF₃, -NR₁₀C(Z)OR₁₀, 5-(R₁₈)-1,2,4-oxadiazol-3-yl or 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl; m' is 1 or 2; n' is 1 to 10; R₉ is hydrogen, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, C₃₋₇ cycloalkyl, C₅₋₇ cycloalkenyl, aryl, arylalkyl, heteroaryl or heteroarylalkyl or R₈ and R₉ may together with the nitrogen to which they are attached form a heterocyclic ring of 5 to 7 members which ring optionally contains an additional heteroatom selected from oxygen, sulfur or

15 NR₁₂; R₁₄ and R₂₄ is each independently selected from hydrogen, alkyl, nitro or cyano; R₁₅ is hydrogen, cyano, C₁₋₄ alkyl, C₃₋₇ cycloalkyl or aryl; R₁₈ and R₁₉ is each independently selected from hydrogen, C₁₋₄ alkyl, substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl or together denote oxygen or sulfur;

20 and R₂₁ is hydrogen, a pharmaceutically acceptable cation, alkyl, cycloalkyl, aryl, arylC₁₋₄alkyl, heteroaryl, heteroarylC₁₋₄alkyl, heterocycl, aroyl, C₁₋₁₀alkoyl.

25

6. A compound as claimed in claim 5 wherein the substituent Y₁ is selected from halogen, C₁₋₅ alkyl and -(CR₁₀R₂₀)_nY₂ wherein Y₂ is -OR₈, -NO₂, -S(O)_m'R₁₁, -SR₈, -S(O)_mNR₈R₉, -NR₈R₉, -O(CR₁₀R₂₀)_nNR₈R₉, -C(O)R₈, -CO₂R₈, -CO₂(CR₁₀R₂₀)_nCONR₈R₉, -CN, -C(Z)NR₈R₉, -NR₁₀S(O)_mR₁₁, -NR₁₀C(Z)R₈, -NR₁₀C(Z)NR₈R₉, -N(OR₂₁)C(Z)NR₈R₉, -C(Z)NR₈OR₉, -NR₁₀C(=NR₁₅)NR₈R₉, -C(=NOR₁₃)NR₈R₉, 5-(R₁₈)-1,2,4-oxadiazol-3-yl or 4-(R₁₂)-5-(R₁₈R₁₉)-4,5-dihydro-1,2,4-oxadiazol-3-yl.

7. A compound as claimed in claim 6 wherein the group Q has one substituent Y₁ which is selected from -(CR₁₀R₂₀)_nY₂ wherein: n is 0,

1, 2 or 3, and Y_2 is $-OR_8$, $-NO_2$, $-S(O)_m'R_{11}$, $-SR_8$, $-S(O)_mNR_8R_9$, $-NR_8R_9$, $-O(CR_{10}R_{20})_nNR_8R_9$, $-C(O)R_8$, $-CO_2R_8$, $-CO_2(CR_{10}R_{20})_nCONR_8R_9$, $-CN$, $-C(Z)NR_8R_9$, $-NR_{10}S(O)_mR_{11}$, $-NR_{10}C(Z)R_8$, $-C(Z)NR_8OR_9$, $-NR_{10}C(Z)NR_8R_9$, $-N(OR_{21})C(Z)NR_8R_9$,

5 $-C(=NOR_{13})NR_8R_9$, $-NR_{10}C(=NR_{15})NR_8R_9$, 5-(R_{18})-1,2,4-oxadizaol-3-yl and 4-(R_{12})-5-($R_{18}R_{19}$)-4,5-dihydro-1,2,4-oxadiazol-3-yl.

8. A compound as claimed in claim 7 wherein, in Y_1 , n is 0 or 1 and Y_2 is $-OH$, $-S(O)_m'R_{11}$, $-SR_8$, $-NR_8R_9$, $-CO_2R_8$, $-S(O)_mNR_8R_9$, $-NR_{10}S(O)_mR_{11}$, 5-(R_{18})-1,2,4-oxadizaol-3-yl or 4-(R_{12})-5-($R_{18}R_{19}$)-4,5-dihydro-1,2,4-oxadiazol-3-yl.

10 9. A compound as claimed in any one of claims 1 to 8 wherein, in R_4 , a 4-phenyl, 4-naphth-1-yl or 5-naphth-2-yl substituent is halogen,

15 $-SR_5$, $-SOR_5$, $-OR_{36}$, $-(CR_{10}R_{20})_mNR_{16}R_{26}$, or $-NR_{10}C(Z)R_8$ and a substituent for another positions of substitution is selected from halogen, $-S(O)_mR_8$, $-OR_8$, $-(CR_{10}R_{20})_mNR_{16}R_{26}$, $-NR_{10}C(Z)R_8$ or $-NR_{10}S(O)_mR_{11}$.

20 10. A compound as claimed in any one of claims 1 to 9 wherein R_1 is 4-pyridyl, 2-alkyl-4-pyridyl or 4-quinolyl; R_2 is hydrogen or methyl; R_3 is phenyl or phenyl substituted, preferably at the 4-position, with a substituent selected from $-(CR_{10}R_{20})_nY_2$ wherein Y_2 is wherein n is 0, 1 2 or 3 and Y_2 is $-OR_8$, $-NO_2$, $-S(O)_m'R_{11}$, $-SR_8$, $-S(O)_mNR_8R_9$, $-NR_8R_9$, $-O(CR_{10}R_{20})_nNR_8R_9$, $-C(O)R_8$, $-CO_2R_8$, $-CO_2(CR_{10}R_{20})_nCONR_8R_9$, $-CN$; $-C(Z)NR_8R_9$, $-C(Z)NR_8OR_9$, $-NR_{10}S(O)_mR_{11}$, $-NR_{10}C(Z)R_8$, $-NR_{10}C(Z)NR_8R_9$, $-C(=NOR_{13})NR_8R_9$, $-NR_{10}C(=CR_{14}R_{24})NR_8R_9$, 5-(R_{18})-1,2,4-oxadizaol-3-yl, 4-(R_{12})-5-($R_{18}R_{19}$)-4,5-dihydro-1,2,4-oxadiazol-3-yl or a 3,5-dimethyl- or 3,5-dibromo-4-hydroxyl grouping; and R_4 is phenyl or phenyl substituted at the 4-position with fluoro and/or substituted at the 3-position with fluoro, chloro, C_{1-4} alkoxy, methane-sulfonamido or acetamido.

30 11. A compound of formula (I) as defined in claim 1 selected from:

35 4-(4-Fluorophenyl)-2-(4-methylthiophenyl)-5-(4-pyridyl)imidazole; 4-(4-Fluorophenyl)-2-(4-ethylthiophenyl)-5-(4-pyridyl)imidazole; 4-(4-Fluorophenyl)-2-(4-methylsulfonylphenyl)-5-(4-pyridyl)imidazole; 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)imidazole;

4-(4-Fluorophenyl)-2-(4-ethylsulfinylphenyl)-5-(4-pyridyl)imidazole;
 4-(3-Chlorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)imidazole;
 2-[4-(N-Methyl-N-benzyl)aminomethylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)imidazole;

5 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylthiophenyl)-imidazole;
 4-(4-Fluorophenyl)-5-[4-(2-methylpyridyl)]-2-(4-methylsulfinylphenyl)-imidazole;
 4-(4-Fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-quinolyl)imidazole;

10 2-[4-(N-Morpholino)methylphenyl]-4-(4-fluorophenyl)-5-(4-pyridyl)-imidazole; or
 a pharmaceutically acceptable salt thereof.

12. A process for preparing a compound of formula (I) as defined in
 15 any one of claims 1 to 11 which process comprises:
 (i) condensing an α -diketone of formula (II):



wherein R_1 and R_4 are as hereinbefore defined, or an equivalent thereof,
 20 with an aldehyde of the formula (III):



wherein R_3 is as hereinbefore defined, or an equivalent thereof, and, if necessary, with ammonia or a source thereof, under imidazole-ring forming conditions;

25 (ii) treating an α -hydroxyketone compound of formula (IIA):



wherein one of R' and R'' is R_1 and the other is R_4 , a suitably protected derivative thereof or the α -hydroxy-oxime or α -haloketone derivative thereof, with an oxidising agent capable of converting said compound 30 into the corresponding α -diketone, in the presence of an aldehyde of formula (III) or an equivalent thereof, and a source of ammonia;

(iii) treating an amidine of formula (IV):

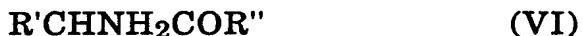


wherein R_2 and R_3 are as hereinbefore defined, or a salt thereof, with 35 a reactive ester of an α -hydroxyketone of formula (IIA) or the corresponding α -haloketone, in an inert solvent, at a moderately elevated temperature and, if necessary, in the presence of a suitable condensation agent;

(iv) treating an iminoether of formula (V):

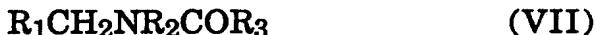


wherein R_3 is as hereinbefore defined and R is C_{1-10} alkyl, aryl or aryl C_{1-4} alkyl, with an α -aminoketone of the formula (VI):



5 wherein one of R' and R'' is R_1 and the other is R_4 in a suitable solvent;

(v) treating the anion of an amide of formula (VII):



wherein R_1 and R_3 are as hereinbefore defined and R_2 is as

10 hereinbefore defined other than hydrogen, with:

(a) a nitrile of the formula (VIII):



wherein R_4 is as hereinbefore defined, or

(b) an excess of an acyl halide, of the formula (IX):



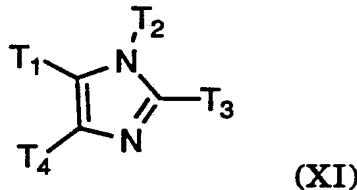
wherein R_4 is as hereinbefore defined and Hal is halogen, or a corresponding anhydride, to give a *bis*-acylated intermediate which is then treated with a source of ammonia;

(vi) treating a compound of formula (X):



wherein R' , R'' and R_4 are as hereinbefore defined and X_c is O or NH, with a source of ammonia, or cyclising the corresponding Schiff's base, formed by treating the compound of formula (X) with an amine R_2NH_2 ;

25 (vii) coupling a suitable derivative of a compound of formula (XI):



wherein: T_2 is a nitrogen protecting group or R_2 , other than hydrogen; and T_1 is hydrogen, T_3 is Q and T_4 is R_4 ; T_1 is R_1 , T_3 is

30 hydrogen and T_4 is R_4 ; or T_1 is R_1 , T_3 is Q and T_4 is hydrogen, in which R_1 , R_2 , R_3 , R_4 and Q are as hereinbefore defined; with: (i) when T_1 is hydrogen, a suitable derivative of the heteroaryl ring R_1H , under ring coupling conditions, to effect coupling of the heteroaryl ring R_1 to the imidazole nucleus at position 5; (ii) when T_3 is hydrogen, a suitable derivative of the aryl or heteroaryl ring QH ,

under ring coupling conditions, to effect coupling of the ring Q to the imidazole nucleus at position 2; or (iii) when T₄ is hydrogen, a suitable derivative of the aryl ring R₄H, under ring coupling conditions, to effect coupling of the aryl ring R₄ to the imidazole nucleus at position 4;

5 (viii) treating a compound of formula (XI), wherein T₁ is hydrogen, with an N-acyl heteroaryl salt, to give an intermediate in which the heteroaryl ring is attached to the imidazole nucleus and is present as a 1,4-dihydro derivative thereof, which intermediate is then 10 subjected to oxidative-deacylation conditions; and

thereafter and if necessary carrying out all or any of the additional steps of removing a protecting group, transforming an initially obtained compound of formula (I) into a further compound of formula (I) or forming a pharmaceutically acceptable salt.

15 13. A compound of formula (I), as defined in any one of claims 1 to 11, or a pharmaceutically acceptable salt thereof, for use in therapy.

20 14. The use of a compound of formula (I), as defined in any one of claims 1 to 11, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for treating a cytokine-mediated disease state.

25 15. A pharmaceutical composition comprising an effective, non-toxic amount of a compound of formula (I), as defined in any one of claims 1 to 11, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier or diluent.

INTERNATIONAL SEARCH REPORT

T/US 93/00674

International Application No

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all)⁶

According to International Patent Classification (IPC) or to both National Classification and IPC
 Int.C1. 5 C07D401/04; C07D403/04; A61K31/44;
 A61K31/505 A61K31/47

II. FIELDS SEARCHED

Minimum Documentation Searched⁷

Classification System	Classification Symbols
Int.C1. 5	C07D ; C07F

Documentation Searched other than Minimum Documentation
 to the Extent that such Documents are Included in the Fields Searched⁸

III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹

Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
X	DE,A,2 221 546 (CIBA-GEIGY A.G.) 16 November 1972 see claims 1,31-33,36,38,43,49,51 see claims 52,61-64 & US,A,3 929 807 cited in the application ---	1-7, 12-15
X	US,A,3 772 441 (JOSEPH GEORGE LOMBARDINO) 13 November 1973 see claims & US,A,3 707 405 cited in the application ---	1-5, 12-15
A	EP,A,0 011 111 (SMITH KLINE CORPORATION) 28 May 1980 see claims -----	1,13-15

* Special categories of cited documents :¹⁰

- "A" document defining the general state of the art which is not considered to be of particular relevance
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"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

1 Date of the Actual Completion of the International Search 14 JUNE 1993	Date of Mailing of this International Search Report 21. 06. 93
International Searching Authority EUROPEAN PATENT OFFICE	Signature of Authorized Officer HENRY J.C.

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9300674
SA 70064

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